

FORM PTO-1390 U S DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTORNEY'S DOCKET NUMBER 4239-58378
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. § 371		U S APPLICATION NO (If known, see 37 C.F.R. § 1.5) 09/807148
INTERNATIONAL APPLICATION NO. PCT/US99/23240	INTERNATIONAL FILING DATE October 5, 1999	PRIORITY DATE CLAIMED October 6, 1998
TITLE OF INVENTION USE OF CALRETICULIN AND CALRETICULIN FRAGMENTS TO INHIBIT ENDOTHELIAL CELL GROWTH AND ANGIOGENESIS, AND SUPPRESS TUMOR GROWTH		
APPLICANT(S) FOR DO/EO/US Giovanna Tosato, Sandra E. Pike, and Lei Yao		
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information		
<ol style="list-style-type: none"> 1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. § 371 2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. § 371. 3. <input type="checkbox"/> This express request to begin national examination procedures (35 U.S.C. § 371(f) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. § 371(b) and PCT Articles 22 and 39(1) 4. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date. 5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. § 371(c)(2)) <ol style="list-style-type: none"> a. <input type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau). b. <input type="checkbox"/> has been transmitted by the International Bureau c. <input checked="" type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US). 6. <input type="checkbox"/> A translation of the International Application into English (35 U.S.C. § 371(c)(2)) 7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. § 371(c)(3)) <ol style="list-style-type: none"> a. <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau) b. <input type="checkbox"/> have been transmitted by the International Bureau. c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. d. <input checked="" type="checkbox"/> have not been made and will not be made 8. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. § 371(c)(3)) 9. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. § 371(c)(4)) 10. <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. § 371(c)(5)) 		
Items 11. to 16. below concern document(s) or information included:		
<ol style="list-style-type: none"> 11. <input checked="" type="checkbox"/> An Information Disclosure Statement under 37 C.F.R. §§ 1.97 and 1.98. 12. <input checked="" type="checkbox"/> Letter to the Official Draftsperson 13. <input checked="" type="checkbox"/> A FIRST preliminary amendment. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment. 14. <input type="checkbox"/> A substitute specification. 15. <input type="checkbox"/> A change of power of attorney and/or address letter. 16. <input checked="" type="checkbox"/> Other items or information: <input type="checkbox"/> Written Opinion. <input checked="" type="checkbox"/> Preliminary Examination Report. <input checked="" type="checkbox"/> International Search Report <input checked="" type="checkbox"/> Copies of References Cited (in Information Disclosure Statement) 		



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U.S. APPLICATION NO. (if known, see 37 C.F.R. § 1.53) 09/807148		INTERNATIONAL APPLICATION NO. PCT/US99/23240		ATTORNEY'S DOCKET NUMBER 4239-58378	
17. <input checked="" type="checkbox"/> The following fees are submitted: BASIC NATIONAL FEE (37 C.F.R. §§ 1.492(a)(1)-(5)): Neither International Preliminary Examination fee (37 C.F.R. § 1.482) nor International Search fee (37 C.F.R. § 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO..... \$1,000.00 International Preliminary Examination fee (37 C.F.R. § 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO..... \$860.00 International Preliminary Examination fee (37 C.F.R. § 1.482) not paid to USPTO but International Search fee (37 C.F.R. § 1.445(a)(2)) paid to USPTO..... \$710.00 International Preliminary Examination fee paid to USPTO (37 C.F.R. § 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4)..... \$690.00 International Preliminary Examination fee paid to USPTO (37 C.F.R. § 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4)..... \$100.00				CALCULATIONS (PTO USE ONLY)	
ENTER APPROPRIATE BASIC FEE AMOUNT =				\$	860.00
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 C.F.R. § 1.492(e)).				\$	0.00
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE		
Total claims	59 - 20 =	39	x \$18.00	\$	702.00
Independent Claims	12 - 3 =	9	x \$80.00	\$	720.00
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$270.00	\$	
TOTAL OF ABOVE CALCULATIONS =				\$	2,282.00
<input type="checkbox"/> Reduction of 1/2 for filing by small entity Small entity status is claimed for this application.				\$	
SUBTOTAL =				\$	2,282.00
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 Months from the earliest claimed priority date (37 C.F.R. §§ 1.492(f)).				\$	
TOTAL NATIONAL FEE =				\$	2,282.00
Fee for recording the enclosed assignment (37 C.F.R. § 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 C.F.R. §§ 3.28, 3.31). \$40.00 per property.				\$	
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b. <input type="checkbox"/> Please charge my Deposit Account No. _____ in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.					
c. <input checked="" type="checkbox"/> The Director is hereby authorized to charge any additional fees that may be required, or credit any overpayment, to Deposit Account No. <u>02-4550</u> . A duplicate copy of this sheet is enclosed.					
d. <input checked="" type="checkbox"/> Please return the enclosed postcard to confirm that the items listed above have been received.					
NOTE: Where an appropriate time limit under 37 C.F.R. § 1.494 or § 1.495 has not been met, a petition to revive (37 C.F.R. § 1.137(a) or (b)) must be filed and granted to restore the application to pending status.					
SEND ALL CORRESPONDENCE TO:				<u>William D Noonan</u> SIGNATURE	
KLARQUIST SPARKMAN CAMPBELL LEIGH & WHINSTON, LLP One World Trade Center, Suite 1600 121 S.W. Salmon Street Portland, OR 97204-2988				William D. Noonan, M.D. NAME	
				<u>43,121</u> REGISTRATION NUMBER	

cc: Docketing

09/807148

EXPRESS MAIL LABEL NO. EL754020413US

DATE OF DEPOSIT: April 5, 2001

PATENT

JCOB Rec'd PCT/PTO 05 APR 2001

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of: Tosato et al.

Application No.: To be assigned

Filed: Herewith

For: USE OF CALRETICULIN AND
CALRETICULIN FRAGMENTS TO INHIBIT
ENDOTHELIAL CELL GROWTH AND
ANGIOGENESIS, AND SUPPRESS TUMOR
GROWTH

Examiner: To be assigned

Date: April 5, 2001

Art Unit: To be assigned

CERTIFICATE OF EXPRESS MAILING

I hereby certify that this paper and the documents referred to as being attached or enclosed herewith are being deposited with the United States Postal Service on April 5, 2001 as Express Mail Label No. EL754020413US in an envelope addressed to: BOX PCT, COMMISSIONER FOR PATENTS, WASHINGTON, D.C. 20231.



William D. Noonan, M.D.
Attorney for Applicant

BOX PCT
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WASHINGTON, D.C. 20231

PRELIMINARY AMENDMENT

Prior to calculating the fees for the above-identified application, please enter the following amendments:

In the claims:

Please add the following claims:

57. (New) The method of claim 1, wherein the therapeutically effective fragment of calreticulin consists essentially of an amino acid sequence selected from the group consisting of:
- (a) the amino acid sequence shown in SEQ ID NO: 5
 - (b) the amino acid sequence shown in SEQ ID NO: 6;
 - (c) the amino acid sequence shown in SEQ ID NO: 8;
 - (d) the amino acid sequence shown in SEQ ID NO: 9; and
 - (e) amino acid sequences comprising fragments and variants of the sequences of (a), (b), (c), and (d), wherein the amino acid sequence inhibits tumor growth.

58. (New) A method of inhibiting radiation induced injury, comprising contacting cells with a pharmaceutical composition comprising at least one protein selected from the group consisting of:

- (a) therapeutically effective fragments of calreticulin;
- (b) therapeutically effective variants of calreticulin; and
- (c) calreticulin.

59. (New) A method of inhibiting chemotherapy induced injury, comprising contacting cells with a pharmaceutical composition comprising at least one protein selected from the group consisting of:

- (a) therapeutically effective fragments of calreticulin;
- (b) therapeutically effective variants of calreticulin; and
- (c) calreticulin.

REMARKS

New claims 57-59 have been added. Support for new claim 57 can be found in the specification on page 9, lines 12-13, and on page 3, lines 23-31. Support for new claims 58 and 59 can be found in the specification on page 39, lines 22-24. No new matter has been added. Applicant respectfully requests examination of the application.

CONCLUSION

If any issues remain before examination of the subject application, the Examiner is respectfully requested to telephone the undersigned Applicants' representative.

Respectfully submitted,

KLARQUIST SPARKMAN CAMPBELL
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**USE OF CALRETICULIN AND CALRETICULIN FRAGMENTS TO INHIBIT
ENDOTHELIAL CELL GROWTH AND ANGIOGENESIS, AND SUPPRESS TUMOR
GROWTH**

5

Field of the Invention

This invention relates to inhibition of endothelial cell growth, angiogenesis and tumor growth.

Background

A. Tumor Growth and Invasion

10 Tumor growth and invasion into normal tissues is dependent upon an adequate blood supply (Folkman, *Ann. N.Y. Acad. Sci.* 401:212-227, 1982; Kerbel, *Bioessays* 13:31-36, 1991). Agents that target tumor blood supply have been shown to prevent or delay tumor formation, and to promote the regression or dormancy of established tumors in preclinical models. Thus, antibodies against the endothelial cell growth factor VEGF (vascular endothelial growth factor), which is produced at high
15 levels by various types of tumors (Dvorak et al., *J. Exp. Med.* 174:1275-1278, 1991), antibodies to VEGF receptor 2, and soluble VEGF receptors all have been shown to reduce tumor growth in experimental animal models (Kendall and Thomas, *Proc. Natl. Acad. Sci. U.S.A.* 90:10705-10709, 1993; Kim et al., *Nature* 362:841-844, 1993; Skobe et al., *Nat. Med.* 3:1222-1227, 1997). Antibodies to the integrin $\alpha_v\beta_3$, which is expressed at high levels by angiogenic blood vessels and permits
20 endothelial cells to interact with components of the extracellular matrix, have been shown to disrupt ongoing angiogenesis on the chick chorioallantoid membrane and produce regression of human tumors transplanted into this site (Cheresh and Spiro, *J. Biol. Chem.* 262:17703-17711, 1987; Brooks et al., *Cell* 79:1157-1164, 1994).

A truncated form of tissue factor targeted to tumor vascular endothelium was demonstrated
25 to initiate formation of intravascular clots and promote the regression of experimental tumors established in mice (Huang et al., *Science* 275:547-550, 1997). Angiostatin, a fragment of plasminogen (O'Reilly et al., *Cell* 79:315-328, 1994), and endostatin, a fragment of collagen XVIII (O'Reilly et al., *Cell* 88:277-285, 1997), are known to inhibit the proliferation of endothelial cells *in vitro* and to suppress neovascularization *in vivo*. Both compounds also inhibit the growth of a variety
30 of tumors in mice, and upon repeated cycles of treatment, promote sustained tumor dormancy without inducing drug resistance (O'Reilly et al., *Nat. Med.* 2:689-692, 1996; Boehm et al., *Nature* 390:404-407, 1997).

Other multifunctional drugs that can also inhibit angiogenesis have displayed antitumor effects. These include Interleukin-12 (Voest et al., *J. Natl. Cancer Inst.* 87:581-586, 1995) the
35 Interferon- γ Inducible Protein-10 (Angiolillo et al., *J. Exp. Med.* 182:155-162, 1995; Strieter et al., *Biochem. Biophys. Res. Commun.* 210:51-57, 1995; Sgadari et al., *Proc. Natl. Acad. Sci. U.S.A.* 93:13791-13796, 1996), the monokine induced by Interferon- γ (Sgadari et al., *Blood* 89:2635-2643, 1997), a fragment of prolactin (Clapp et al., *Endocrinology* 133:1292-1299, 1993), synthetic analogues of fumagillin (Ingber et al., *Nature* 348:555-557, 1990), thalidomide (D'Amato et al., *Proc*

Natl Acad Sci U.S.A. 91:4082-4085, 1994), Platelet Factor-4 (Maione et al., *Science* 247:77-79, 1990), and thrombospondin (Good et al., *Proc Natl Acad Sci U.S.A.* 87:6624-6628, 1990; Weinstat-Saslow et al., *Cancer Res.* 54:6504-6511, 1994).

5 **B. Calreticulin**

Calreticulin was first identified in skeletal muscle sarcoplasmic reticulum. (Ostwald and MacLennan, *J. Biol. Chem.* 249:974-979, 1974). Fifteen years later it was cloned and the N-terminus was sequenced. This led to the discovery that several groups had independently identified the molecule and had given it different names, including, "high-affinity Ca^{2+} ", "calregulin", "CRP55"
10 and "calsequestrin-like protein" (Ostwald and MacLennan, *J. Biol. Chem.* 249:974-979, 1974; Waisman et al., *J. Biol. Chem.* 260:1652-1660, 1985; Macer, D. R. J. & Koch, G. L. E. *J. Cell. Sci.* 91:61-70, 1988; Damiani et al., *Biochem Biophys Res Commun* 165:973-980, 1989; Treves et al., *Biochem. J.* 271:473-480, 1990). Each of these groups identified calreticulin through different means, but all identified its ability to bind Ca^{2+} .

15 Although most studies have indicated that calreticulin resides predominantly within the lumen of the endoplasmic reticulum, calreticulin may also be found in other cellular compartments. For example, calreticulin was detected on the plasma membranes of lymphoblastoid cells (Newkirk and Tsoukas, *J. Autoimmun.* 5:511-525, 1992) and epidermal keratinocyte lines (Kawashima, et al., *Dermatology* 189 Suppl. 1:6-10, 1994). It was proposed to represent, or to be closely related in
20 structure, to the C1q receptor found on endothelial cells, B cells, T cells and other cells (Chen et al., *J. Immunol.* 153:1430-1440, 1994). Calreticulin is also a constituent of lytic granules contained in cytotoxic T and NK cells from which it is released during cell lysis (Dupuis et al., *J. Exp. Med.* 177:1-7, 1993), and has been purified from the culture supernatant of several cell types (Booth and Koch, *Cell* 59:729-737, 1989; Eggleton et al., *Clin. Immunol. Immunopathol.* 72:405-409, 1994) and
25 from normal human plasma (Sueyoshi et al., *Thromb. Res.* 63:569-575, 1991). Several observations support the notion that calreticulin can also be a target for autoimmune responses (Lux et al., *J. Clin. Invest.* 89:1945-1951, 1992; Meilof et al., *J. Immunol.* 151:5800-5809, 1993).

Since the initial identification and cloning, the structure of calreticulin has been characterized. Mammalian calreticulin is a 417 amino acid peptide from which the 17 N-terminal
30 amino acids are cleaved upon translocation to the lumen of the endoplasmic reticulum (Smith and Koch, *Embo. J.* 8:3581-3586, 1989). In addition to being found in the lumen of the endoplasmic reticulum, calreticulin has been found in the cytoplasm, in the nucleus of some cells, and in the extracellular matrix (Michalak et al., *Biochem. J.* 285:681-692, 1992). Further studies revealed that calreticulin has three distinct domains, the N-terminal domain, a middle domain and the C-terminal
35 domain.

The mature calreticulin is composed of an N-terminal domain consisting of 180 amino acids that are highly conserved. Proposed three-dimensional models indicate that the domain contains eight anti-parallel β -strands. Furthermore, the N-terminal domain has been found to bind a number

of molecules including the alpha subunit of integrin, Zn^{2+} , and the DNA binding domain of steroid receptors (Nash et al., *Mol. Cellular Biochem.* 135:71-78, 1994).

The middle domain of calreticulin stretches from amino acid 180 to amino acid 280. It is proline rich and has also been termed the P-domain. This domain has been found to have a high
 5 affinity for Ca^{2+} and contains a nuclear localization signal (Baksh and Michalak, *J. Biol. Chem.* 266:21458-21465, 1991).

Following the P-domain is the C-domain. This last domain is highly acidic and contains an endoplasmic reticulum retention signal. The C-domain binds to Factor IX, Factor X, and prothrombin (See U.S. Patent No 5,426,097, to Stern et al.).

10 Calreticulin has also been found to be useful in wound healing (See U.S. Patent No 5,591,716, to Siebert et al.).

Summary

Calreticulin has been actively studied since 1972. Some of the studies that have been done
 15 have focused on understanding the structure of calreticulin while others have focused on understanding the physiological role of the molecule.

The present invention stems from the discovery that calreticulin has three previously uncharacterized biological activities. First, calreticulin is shown to inhibit endothelial cell growth, while having little or no effect on the growth of non-endothelial cells. Second, calreticulin is shown
 20 to inhibit angiogenesis. Third, calreticulin is shown to inhibit tumor growth, including the growth of Burkitt lymphoma, breast adenocarcinomas, colon carcinomas, lung carcinomas, melanoma, rhabdomyosarcoma, promyelomonocytic lymphoma, Wilms tumor, and neuroblastoma tumors.

It has also been discovered that certain fragments of calreticulin share these activities. These fragments include calreticulin lacking the N-terminal 1-120 amino acids (Seq. I.D. No. 9), the N-terminal domain (Seq. I.D. No. 4), as well as fragments of the N-terminal domain of calreticulin.
 25 Examples of these fragments include the recombinantly produced 180 amino acid N-terminal domain of calreticulin (Seq. I.D. No. 4), which has been given the name "vasostatin", Δ 120 calreticulin (calreticulin missing the N-terminal 1-120 amino acids; Seq. I.D. No. 9), as well as a recombinantly produced 61 amino acid fragment (a.a. 120-180; Seq. I.D. No. 5). Other biologically active
 30 fragments of calreticulin include a synthetic 49 amino acid fragment (a.a. 132-180; Seq. I.D. No. 6) and a synthetic 60 amino acid fragment (a.a. 121-180; Seq. I.D. No. 8).

Additionally, the activity of the above described molecules does not stem from their ability to bind to the sequence $KXaa_1FFXaa_2R$ (Seq. I.D. No. 11; Xaa_1 represents either G, A, or V, and Xaa_2 represents K or R) or the sequence $KXaa_1GFFKR$ (Seq. I.D. No. 10; Xaa_1 represents either I, L,
 35 G, A, or C) that are known to bind calreticulin.

Calreticulin and the described fragments of calreticulin are useful among other things for inhibiting endothelial cell growth, for the treatment of subjects having solid tumors with associated neovascularization, and for the treatment of other diseases where angiogenesis is a factor. In particular, the present invention may be especially useful for treating rheumatoid arthritis,

autoimmune diseases, rheumatic diseases, and certain ocular neovascular diseases, such as, macular degeneration, diabetic retinopathy, and retrolental fibroplasia.

Pharmaceutical compositions comprising calreticulin and the therapeutically active fragments and variants of calreticulin are provided by the invention. In particular embodiments, the biologically active fragments are those fragments having the sequences shown in Seq. I.D. Nos. 4, 5, 6, 8, and 9.

In other aspects, the invention provides a method of inhibiting endothelial cell growth by contacting endothelial cells with a pharmaceutical composition that comprises at least one protein selected from the group consisting of calreticulin, and therapeutically effective fragments of calreticulin. The invention also provides a method of inhibiting angiogenesis in a subject, by administering to the subject an effective amount of a pharmaceutical composition comprising at least one protein selected from the group consisting of calreticulin and therapeutically effective fragments and variants of calreticulin.

With respect to the anti-tumor activities of the molecules provided, another aspect of the invention is a method of inhibiting tumor angiogenesis and growth by contacting tumor cells with an effective amount of a pharmaceutical composition comprising at least one protein selected from the group consisting of calreticulin and therapeutically effective fragments and variants of calreticulin. The anti-tumor activity may also be utilized in a method of inhibiting tumor growth in a subject. That method comprises administering to the subject an effective amount of a pharmaceutical composition comprising at least one protein selected from the group consisting of calreticulin and therapeutically effective fragments and variants of calreticulin.

Among the compositions provided by the invention is a protein with an amino acid sequence selected from the group consisting of therapeutically effective fragments of vasostatin and variants of calreticulin that do not bind to the sequence shown in Seq. I.D. No. 11 and/or Seq. I.D. No. 10, but yet display one of the three biological activities described above.

The invention also provides methods of identifying therapeutically effective variants and fragments of calreticulin. These methods involve contacting a sample suspected of containing therapeutically effective variants or fragments with the sequence KXaa₁FFXaa₂R (Seq. I.D. No. 11), and/or the sequence KXaa₁GFFKR (Seq. I.D. No. 10) detecting the portion of the sample that does not bind to the sequence. The unbound portion of the sample is then tested for one of the three biological activities described above. Accordingly, the invention also provides the therapeutically effective variants and fragments of calreticulin identified by the method, as well as cells containing recombinant nucleic acid molecules that encode the therapeutically effective variants and fragments.

These and other aspects of the invention are explained in more detail in the following sections.

Brief Description of the Drawings

Figure 1 is a graph showing the inhibition of endothelial cell proliferation by MBP-calreticulin (Seq. I.D. No. 2). Fetal bovine heart endothelial cells (800 cells/well) were incubated for

5 days either in medium alone or medium supplemented with bFGF (25 ng/ml), with or without recombinant purified MBP-calreticulin (Seq. I.D. No. 3) or MBP (both at 1 μ g/ml). The results of 16 experiments are expressed as mean cpm (\pm SD).

Figure 2 is a graph showing the dose dependency of MBP-vasostatin (Seq. I.D. No. 4) inhibition.

Figure 3 shows the inhibition of endothelial cell proliferation by Δ 120 calreticulin (Seq. I.D. No. 9). Fetal bovine heart endothelial cells (800 cells/well) were cultured for 5 days in medium alone or medium supplemented with bFGF (25 ng/ml). Recombinant purified MBP-calreticulin (Seq. I.D. No. 3), MBP- Δ 120 calreticulin (Seq. I.D. No. 9), or MBP were added to bFGF-supplemented cultures. Proliferation was measured by 3 H-thymidine incorporation during the final 20-23 hours of culture. The results reflect the mean of triplicate cultures; SDs within 12% of the mean.

Figure 4 is a graph showing the inhibition of endothelial cell proliferation by the 61 amino acid calreticulin fragment MBP-120-180 calreticulin (Seq. I.D. No. 5). Fetal bovine heart endothelial cells (800 cells/well) were incubated for 5 days either in medium alone or medium supplemented with bFGF (15 ng/ml), with or without the recombinant purified 61 amino acid fragment (Seq. I.D. No. 5) (both at 1 μ g/ml). The results of 8 experiments are expressed as mean cpm (\pm SD).

Figure 5 shows a graph comparing endothelial cell growth inhibition by MBP calreticulin (Seq. I.D. No. 3), MBP- Δ 120 calreticulin (Seq. I.D. No. 9), MBP-vasostatin (Seq. I.D. No. 4), and MBP-120-180 calreticulin fragment (Seq. I.D. No. 5). Fetal bovine heart endothelial cells (800 cells/well) were cultured for 5 days in medium alone or medium supplemented with bFGF (15 ng/ml). Recombinant purified fusion proteins were added to culture at 0.4-32 nM concentrations to bFGF-supplemented cultures. Proliferation was measured by 3 H thymidine incorporation during the final 20-23 hours of culture. The results reflect the mean of triplicate cultures; SDs within 15% of the mean. The mean response of endothelial cells was 2,217 c.p.m. when cultured in medium alone, and 23,377 c.p.m. when cultured with bFGF alone.

Figures 6A, 6B, 6C and 6D are graphs showing the inhibition of tumor growth by vasostatin (Seq. I.D. No. 4) and calreticulin (Seq. I.D. No. 3). Burkitt lymphoma cells (CA46 cell line, 1×10^7 cells) were inoculated subcutaneously (s.c.) into BALB/c athymic mice, 6 weeks of age. Beginning on the day of cell inoculation and continuing thereafter daily, 6 days/week, mice were inoculated s.c. with either control buffer (open symbols) or test protein (closed symbols). The results reflect the % mice with tumor as a function of time. **A.** 12 mice were inoculated with control purified GST protein (Seq. I.D. No. 2; 20 μ g/day x 14 days), and 13 mice were inoculated with purified GST-calreticulin (Seq. I.D. No. 2; 60 μ g/day x 14 days). **B.** 8 mice were inoculated with control purified MBP protein (20 μ g/day x 14 days), and 9 mice were inoculated with purified MBP-vasostatin (Seq. I.D. No. 4; 30 μ g/day x 14 days). **C.** 12 mice were inoculated with control purified MBP (40 μ g/day x 18 days), and 12 mice were inoculated with MBP-vasostatin (Seq. I.D. No. 4; 60 μ g/day x 18 days); all mice with tumor (12 treated with MBP and 4 treated with MBP-vasostatin) were killed on day 18. The remaining mice were observed up to day 60. **D.** 6 mice were treated

with formulation buffer alone (0.1 ml/day x 22 days, open circles), 5 mice were treated with 20 µg/day x 22 days purified MBP-vasostatin (Seq. I.D. No. 4; diamonds), and 9 mice were treated with 100 µg/day x 22 days purified MBP-vasostatin (Seq. I.D. No. 4; squares).

Figure 7A and 7B are graphs showing the inhibition of established Burkitt tumor growth by vasostatin. Burkitt lymphoma cells (CA46 cell line, 1×10^7 cells) were inoculated s.c. into BALB/c athymic mice, 6 weeks of age. **A.** After a tumor appeared (at least 160 mm² in size), 9 mice were treated with MBP-vasostatin (Seq. I.D. No. 4; 200 µg/day, 6 days/week, 100 µl/dose x 46 days) and 10 mice were treated with formulation buffer alone (100 µl x 46 days). Tumor size was recorded. Formulation buffer (open squares); MBP-vasostatin (Seq. I.D. No. 4; closed circles) **B.** After a tumor appeared (at least 160 mm² in size), 12 mice were treated with MBP-vasostatin (Seq. I.D. No. 4; 200 µg/twice/day, 5 days/week and once/day 2 days/week, 100 µl/dose x 26 days) and 8 mice were treated with formulation buffer alone (100 µl x 26 days). The % mice with tumor was determined. Formulation buffer (open squares); MBP vasostatin (Seq. I.D. No. 4; closed squares).

Figure 8 is a graph showing the inhibition of human colon carcinoma growth by vasostatin (Seq. I.D. No. 4). Mice (BALB/c athymic mice, 6 weeks of age) were inoculated s.c. with the human colon carcinoma cell line (SW-480 from ATCC, 6×10^6 cells/mouse in 0.2 ml RPMI medium). After a tumor appeared (at least 130 mm² in size), 12 mice were treated with MBP-vasostatin (Seq. I.D. No. 4; 100 µg/day, 6 days/week, 100 µl/dose, x 36 days; closed circles) and 10 mice were inoculated with formulation buffer (100 µl x 36 days; open circles). Tumor size was estimated as the product of two-dimensional caliper measurements.

Figures 9A and 9B are graphs showing the inhibition of human neuroblastoma tumor growth. Mice (BALB/c athymic mice, 6 weeks of age) were inoculated s.c. with the human neuroblastoma cell line (SK-N-MC from ATCC, 8.5×10^6 cells/mouse in 0.2 ml RPMI medium). Beginning on the day of cell inoculation and continuing thereafter daily, 6 days/week, 12 mice were inoculated s.c. with control MBP (20 µg/day) and 13 mice were inoculated with MBP-vasostatin (Seq. I.D. No. 4; 30 µg/day). On day 30, 8 of the 12 mice that had been injected with control MBP (mean tumor weight 1.74 g) and 5 of the 13 mice injected with MBP-vasostatin (Seq. I.D. No. 4; mean tumor weight 1.78 g) were sacrificed. Treatment was continued in the remaining mice until day 55 at which time all animals were sacrificed. One of the mice treated with MBP-vasostatin (Seq. I.D. No. 4) did not develop a tumor and since day 55 was observed untreated; as of day 220 no tumor has developed. On day 55, all animals with tumor were sacrificed and tumor weight measured. There were 4 animals in the control group (MBP) and 7 animals in the treatment group (MBP-vasostatin). **A.** Tumor growth curves. **B.** Comparison of tumor weights in animals sacrificed on day 55.

Figure 10 is a graph showing the inhibition of human breast adenocarcinoma tumor growth by vasostatin (Seq. I.D. No. 4). Mice (BALB/c athymic mice, 6 weeks of age) were inoculated s.c. with the human breast adenocarcinoma cell line (MDA-MB231 from ATCC, 7.5×10^6 cells/mouse in 0.2 ml RPMI medium). Beginning on the day of cell inoculation and continuing thereafter daily, 5 days/week, for 18 days, 8 mice were inoculated s.c. with control formulation buffer (100 µl/injection/twice/day for 5 days/week and once/day for 2 days/week) and 8 mice were inoculated

with MBP-vasostatin (Seq. I.D. No. 4; 200 µg/twice/day for 5 days/week and once/day for 2 days/week). Tumor size was estimated as the product of two-dimensional caliper measurements.

Figure 11A and 11B are graphs showing the inhibition of murine lymphoma tumor growth by MBP-vasostatin (Seq. I.D. No. 4). Mice (BALB/c athymic mice, 6 weeks of age) were inoculated s.c. with the murine B-cell lymphoma cell line (Raw 8.1 from ATCC, 2.5×10^6 cells/mouse in 0.2 ml RPMI medium). Beginning on the day of cell inoculation and continuing thereafter daily, 7 days/week, 9 mice were inoculated s.c. with control formulation buffer (100 µl/day) and 9 mice were inoculated with MBP-vasostatin (Seq. I.D. No. 4; 200 µg/day). Treatment was continued until all mice developed a tumor measuring at least 0.5 cm² on day 15. A. Tumor size, estimated as the product of two-dimensional caliper measurements, was recorded for control and vasostatin-treated mice. B. Tumor weight in control and vasostatin treated mice measured on day 15.

Figure 12 is a graph showing the inhibition of Wilms tumor growth by MBP-vasostatin (Seq. I.D. No. 4). Athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (12×10^6 cell/mouse) with the human Wilms tumor cell line (SK-NP-1). One day after cell injection, and continuing daily thereafter for 42 days, the mice received either control formulation buffer alone or MBP-vasostatin (Seq. I.D. No. 4; 100 µg/day, 6 days/week). Tumor size was estimated as the product of two dimensional caliper measurement.

Figure 13 is a graph showing the inhibition of rhabdomyosarcoma growth by MBP-vasostatin (Seq. I.D. No. 4). Groups of athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (8×10^6 cell/mouse) s.c. with the human rhabdomyosarcoma cell line A-204 tumor cell line (SK-NP-1). One day after cell injection, and continuing daily thereafter for 28 days, the mice received either control formulation buffer alone or MBP-vasostatin (Seq. I.D. No. 4; 100 µg/day, 6 days/week). Tumor size was estimated as the product of two dimensional caliper measurement.

Figure 14 is a bar graph showing the inhibition of myelomonocytic lymphoma growth by MBP-vasostatin (Seq. I.D. No. 4). Groups of athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (10×10^6 cell/mouse) with the human HL60 tumor cell line. One day after cell injection, and continuing daily thereafter for 24 days, the mice received either control formulation buffer alone or MBP-vasostatin (Seq. I.D. No. 4; 200 µg/day, 6 days/week). The weight of each tumor was measured in grams after tumors were removed from the animals.

Figure 15 is a graph showing the inhibition of human melanoma tumor growth by MBP-vasostatin (Seq. I.D. No. 4). Athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (7×10^6 cell/mouse) with the human melanoma cell line (A-375). Five days after cell injection, and continuing daily thereafter for 20 days, the mice received either control formulation buffer alone or MBP-vasostatin (Seq. I.D. No. 4; 200 µg/day, 6 days/week). Tumor size was estimated as the product of two dimensional caliper measurement.

Figure 16 is a graph showing the inhibition of human Burkitt lymphoma growth by the calreticulin fragment encompassing amino acids 120-180 (Seq. I.D. No. 5). Athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (4×10^6 cell/mouse) with the human Burkitt lymphoma cell line CA46. Beginning on the day of cell inoculation and continuing daily thereafter 6 days/week the mice received either control recombinant purified MBP (65 $\mu\text{g/day}$, 6 days/week), MBP-vasostatin (Seq. I.D. No. 4; 100 $\mu\text{g/day}$, 6 days/week), or MBP-120-180 calreticulin (Seq. I.D. No. 5; 75 $\mu\text{g/day}$, 6 days/week). Mice were injected for 42 days. Tumor size was estimated as the product of two dimensional caliper measurement.

Figures 17A, 17B, 17C, and 17D are graphs showing the inhibition of human Burkitt lymphoma growth by a combination treatment of vasostatin (Seq. I.D. No. 4) and Interleukin 12 (IL-12). Athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (4×10^6 cell/mouse) with the human Burkitt lymphoma cell line CA46. **A and B.** 5 days after cell inoculation the established tumors were treated with daily s.c. inoculations (6 days/week) of either formulation buffer alone (closed circles), MBP-vasostatin alone (Seq. I.D. No. 4; 100 $\mu\text{g/day}$; open squares), mL-12 alone (200 ng/day for 8 days followed by 100 ng/day; open circles), or the combination of MBP-vasostatin (Seq. I.D. No. 4) and mL-12 (same dose used as single agents; open triangles). All mice were sacrificed after 15 days of treatment. **C and D.** 1 day after cell inoculation, s.c. treatment with MBP-vasostatin (Seq. I.D. No. 4; 100 $\mu\text{g/day}$; 6 days/week) or formulation buffer was started; s.c. treatment with IL-12 (100 ng/mouse; 6 days/week) was started 2 days later in mice that had received buffer alone or MBP-vasostatin (Seq. I.D. No. 4). Therefore, four groups of mice were tested: Mice receiving buffer alone (closed circles); MBP-vasostatin alone (open squares); IL-12 alone (open circles); and MBP-vasostatin plus IL-12 (open triangles). The mice received 7 days treatment with MBP-vasostatin (Seq. I.D. No. 4) and 5 days treatment with IL-12, and were observed untreated for 14 days at which time all mice were sacrificed. Tumor size was estimated as the product of two-dimensional caliper measurement.

Figure 18 is a Western blot analysis of fractions obtained through affinity purification over a column containing KXaa₁FFXaa₂R (Seq. I.D. No. 11) and KXaa₁GFFKR (Seq. I.D. No. 10). Purified supernatants from the VDS-0 cell line containing full length calreticulin and N-terminal fragments of calreticulin were affinity purified over a mixture of peptides KLGFFKR (Seq. I.D. No. 13), KAFFKR (Seq. I.D. No. 33) and KVFFKR (Seq. I.D. No. 32) coupled to CNBr-activated Sepharose. Starting material (lane 1), unbound material (lanes 2, 3) and bound material (lanes 4, 5) were separated by SDS-PAGE, transferred to nitrocellulose, and subsequently stained with a rabbit antiserum against human calreticulin.

Sequence Listing

The nucleic and amino acid sequences listed in the accompanying sequence listing are shown using standard letter abbreviations for nucleotide bases, and three letter code for amino

acids. Only one strand of each nucleic acid sequence is shown, but the complementary strand is understood as included by any reference to the displayed strand.

5 Seq. I.D. No. 1 shows the nucleic acid sequence for the open reading frame of human calreticulin.

Seq. I.D. No. 2 shows the amino acid sequence of human calreticulin.

Seq. I.D. No. 3 shows the amino acid sequence of human calreticulin without the 17 N-terminal amino acids (References throughout the text to amino acid numbers are keyed to this sequence).

10 Seq. I.D. No. 4 shows the amino acid sequence of the N-terminal 180 amino acids of human calreticulin ("vasostatin").

Seq. I.D. No. 5 shows the amino acid sequence of the recombinant 61 amino acid fragment of human calreticulin. This sequence corresponds to amino acids number 120-180 of the human calreticulin sequence.

15 Seq. I.D. No. 6 shows the amino acid sequence of the synthetic 49 amino acid fragment of human calreticulin. This sequence corresponds to amino acids numbers 132-180 of the human calreticulin sequence.

Seq. I.D. No. 7 shows the cDNA sequence of human calreticulin.

20 Seq. I.D. No. 8 shows the amino acid sequence of the synthetic 60 amino acid fragment of human calreticulin. This sequence corresponds to amino acids numbers 121-180 of the human calreticulin sequence.

Seq. I.D. No. 9 shows the amino acid sequence of Δ 120 calreticulin (calreticulin (Seq. I.D. No. 3) missing the N-terminal 1-120 amino acids).

25 Seq. I.D. No. 10 shows the integrin consensus amino acid sequence

Seq. I.D. No. 11 shows the steroid nuclear receptor consensus amino acid sequence.

Seq. I.D. No. 12 shows the amino acid sequence of a portion of an integrin.

Seq. I.D. No. 13 shows the amino acid sequence of a portion of an integrin.

Seq. I.D. No. 14 shows the amino acid sequence of a portion of an integrin.

Seq. I.D. No. 15 shows the amino acid sequence of a portion of an integrin.

30 Seq. I.D. No. 16 shows the amino acid sequence of a portion of an integrin.

Seq. I.D. No. 17 shows the amino acid sequence of a portion of an integrin.

Seq. I.D. No. 18 shows the amino acid sequence of a portion of an integrin.

Seq. I.D. No. 19 shows the amino acid sequence of a portion of an integrin.

Seq. I.D. No. 20 shows the amino acid sequence of a portion of an integrin.

35 Seq. I.D. No. 21 shows the amino acid sequence of a portion of a steroid nuclear receptor.

Seq. I.D. No. 22 shows the amino acid sequence of a portion of a steroid nuclear receptor.

Seq. I.D. No. 23 shows the amino acid sequence of a portion of a steroid nuclear receptor.

Seq. I.D. No. 24 shows the amino acid sequence of a portion of a steroid nuclear receptor.

Seq. I.D. No. 25 shows the amino acid sequence of a portion of a steroid nuclear receptor.

Seq. I.D. No. 26 shows the amino acid sequence of a portion of a steroid nuclear receptor.
 Seq. I.D. No. 27 shows the amino acid sequence of a portion of a steroid nuclear receptor.
 Seq. I.D. No. 28 shows the amino acid sequence of a portion of a steroid nuclear receptor.
 Seq. I.D. No. 29 shows the amino acid sequence of a portion of a steroid nuclear receptor.
 Seq. I.D. No. 30 shows the amino acid sequence of a portion of a steroid nuclear receptor.
 Seq. I.D. No. 31 shows the amino acid sequence of a portion of a steroid nuclear receptor.
 Seq. I.D. No. 32 shows the amino acid sequence of a portion of the glucocorticoid receptor.
 Seq. I.D. No. 33 shows the amino acid sequence of a portion of the estrogen receptor.
 Seq. I.D. No. 34 shows the amino acid sequence of a portion of the thyroid receptor.
 Seq. I.D. No. 35 shows the amino acid sequence of a portion of the retinoic acid receptor.

Detailed Description

I. Definitions

Unless otherwise noted, technical terms are used according to conventional usage.

Definitions of common terms in molecular biology may be found in Benjamin Lewin, *Genes V*, published by Oxford University Press, 1994 (ISBN 0-19-854287-9); Kendrew et al. (eds.), *The Encyclopedia of Molecular Biology*, published by Blackwell Science Ltd., 1994 (ISBN 0-632-02182-9); and Robert A. Meyers (ed.), *Molecular Biology and Biotechnology: a Comprehensive Desk Reference*, published by VCH Publishers, Inc., 1995 (ISBN 1-56081-569-8).

Calreticulin: Calreticulin is a calcium binding protein that is found in many animals, and is highly conserved across species. The open reading frame of the prototypical human calreticulin is shown in Seq. I.D. No. 1, while the sequence of the human calreticulin protein is shown in Seq. I.D. No. 2. The present invention is founded on the discovery that calreticulin, and particular fragments of calreticulin, possess certain biological activities ("calreticulin activities"). Specifically, these activities include (1) the inhibition of endothelial cell growth; (2) the inhibition of angiogenesis; and (3) the inhibition of tumor growth.

Each of these activities may be separately assayed by methods described herein. The ability of calreticulin or a fragment of this protein to perform these activities may be beneficial in a number of applications, including clinical applications such as tumor therapy and treatment of diseases with abnormal or excessive angiogenesis.

While the amino acid sequence of the prototypical human calreticulin is shown in Seq. I.D. No. 2, one of skill in the art will appreciate that variations in this amino acid sequence (such as amino acid deletions, additions or substitutions) may be made without substantially affecting the activities of the protein (or fragments of the protein) discussed above. Thus, reference herein to the term "calreticulin" encompasses both the protein having the amino acid sequence shown in Seq. I.D. No. 2, as well as amino acid sequences that are based on this sequence but which include one or more sequence variants. Such sequence variants may also be defined in the degree of amino acid sequence identity that they share with the amino acid sequence shown in Seq. I.D. No. 2. Typically, calreticulin sequence variants will share at least 80% sequence identity with the sequence shown in

Seq. I.D. No. 2. More highly conserved variants will share at least 90% or at least 95% sequence identity with the Seq. I.D. No. 2 sequence. In addition to sharing sequence identity with the prototypical calreticulin protein sequence, such sequence variants possess at least one of the three biological activities noted above.

5 **Therapeutically effective fragments and variants of calreticulin:** It is shown herein that not only does calreticulin possess the specified biological activities (inhibiting endothelial cells, angiogenesis and tumor growth), but that such activities are also found in certain peptide fragments of calreticulin. For example, it is shown that the 180 amino acid N-terminal domain of calreticulin (Seq. I.D. No. 4), which is hereinafter termed "vasostatin," possesses these activities, as do the
10 synthetically produced 49 (Seq. I.D. No. 6) and 60 (Seq. I.D. No. 8) amino acid fragments, as well as the recombinantly produced 61 amino acid fragment (Seq. I.D. No. 5) and Δ 120 calreticulin (Seq. I.D. No. 9). Furthermore, it is shown that the activity of these fragments does not stem from their ability to bind to the amino acid sequence motif KXaa₁FFXaa₂R (Seq. I.D. No. 11), that is found in the a family of steroid receptors (glucocorticoid, mineralcorticoid, progesterone and androgen
15 receptors: KVFFKR (Seq. I.D. No. 32); estrogen receptor: KAFFKR (Seq. I.D. No. 33); thyroid hormone receptor: KSFFRR (Seq. I.D. No. 34); and retinoic acid receptor: KGFFRR (Seq. I.D. No. 35).

 Furthermore, it is shown that the activity of these fragments does not stem from their ability to bind to the amino acid sequence motif found in the alpha subunit of integrins. The consensus
20 sequence representing the alpha subunit of integrin is KXaa₁GFFKR (Seq. I.D. No. 10).

 Hence, the terms "therapeutically effective fragment of calreticulin" or "therapeutically effective variant of calreticulin" includes any calreticulin fragment or variant that, at a minimum, possesses one of the three biological activities noted above. For example, the 61 amino acid calreticulin fragment (Seq. I.D. No. 5) is a therapeutically effective fragment of calreticulin since it
25 possesses the ability to inhibit endothelial cell growth. Whether a given calreticulin fragment or variant possesses one or more of these biological activities can be readily determined by the assays described herein. For example, the ability to inhibit endothelial cell growth can readily be determined for any given fragment of calreticulin using the simple *in vitro* assay described below.

Oligonucleotide: A linear polynucleotide sequence of up to about 100 nucleotide bases in
30 length.

Vector: A nucleic acid molecule as introduced into a host cell, thereby producing a transformed host cell. A vector may include nucleic acid sequences that permit it to replicate in a host cell, such as an origin of replication. A vector may also include one or more selectable marker genes and other genetic elements known in the art.

35 **Transformed:** A transformed cell is a cell into which has been introduced a nucleic acid molecule by molecular biology techniques. As used herein, the term transformation encompasses all techniques by which a nucleic acid molecule might be introduced into such a cell, including transfection with viral vectors, transformation with plasmid vectors, and introduction of naked DNA by electroporation, lipofection, and particle gun acceleration.

Isolated: An "isolated" biological component (such as a nucleic acid or protein or organelle) has been substantially separated or purified away from other biological components in the cell of the organism in which the component naturally occurs, i.e., other chromosomal and extra-chromosomal DNA and RNA, proteins and organelles. Nucleic acids and proteins that have been
5 "isolated" include nucleic acids and proteins purified by standard purification methods. The term also embraces nucleic acids and proteins prepared by recombinant expression in a host cell as well as chemically synthesized nucleic acids.

Sequence identity: the similarity between amino acid sequences is expressed in terms of the similarity between the sequences, otherwise referred to as sequence identity. Sequence identity is
10 frequently measured in terms of percentage identity (or similarity or homology); the higher the percentage, the more similar the two sequences are. Homologs or variants of calreticulin will possess a relatively high degree of sequence identity when aligned using standard methods.

Methods of alignment of sequences for comparison are well known in the art. Various programs and alignment algorithms are described in: Smith and Waterman, *Adv. Appl. Math.* 2:482,
15 1981); Needleman and Wunsch, *J. Mol. Biol.* 48:443, 1970; Pearson and Lipman, *Proc. Natl. Acad. Sci. U.S.A.* 85:2444, 1988; Higgins and Sharp, *Gene* 73:237-244, 1988; Higgins and Sharp, *CABIOS* 5:151-153, 1989; Corpet et al., *Nucleic Acids Research* 16:10881-10890, 1988; and Pearson and Lipman, *Proc. Natl. Acad. Sci. U.S.A.* 85:2444, 1988. Altschul et al., *Nature Genet.*, 6:119-129, 1994, presents a detailed consideration of sequence alignment methods and homology calculations.

20 The NCBI Basic Local Alignment Search Tool (BLAST) (Altschul et al., *J. Mol. Biol.*, 215:403-410, 1990.) is available from several sources, including the National Center for Biotechnology Information (NCBI, Bethesda, MD) and on the Internet, for use in connection with the sequence analysis programs blastp, blastn, blastx, tblastn and tblastx. It can be accessed at <http://www.ncbi.nlm.nih.gov/BLAST/>. A description of how to determine sequence identity using this
25 program is available at http://www.ncbi.nlm.nih.gov/BLAST/blast_help.html.

Homologs and variants of calreticulin are typically characterized by possession of at least 50% sequence identity counted over the full length alignment with the amino acid sequence of calreticulin using the NCBI Blast 2.0, gapped blastp set to default parameters. For comparisons of amino acid sequences of greater than about 30 amino acids, the Blast 2 sequences function is employed using the
30 default BLOSUM62 matrix set to default parameters, (gap existence cost of 11, and a per residue gap cost of 1). When aligning short peptides (fewer than around 30 amino acids), the alignment should be performed using the Blast 2 sequences function, employing the PAM30 matrix set to default parameters (open gap 9, extension gap 1 penalties). Proteins with even greater similarity to the reference sequences will show increasing percentage identities when assessed by this method, such as at least 60%, at least
35 70%, at least 75%, at least 80%, at least 90% or at least 95% or 98% sequence identity. When less than the entire sequence is being compared for sequence identity, homologs and variants will typically possess at least 75% sequence identity over short windows of 10-20 amino acids, and may possess sequence identities of at least 85% or at least 90% or 95% depending on their similarity to the reference sequence. Methods for determining sequence identity over such short windows are described at

http://www.ncbi.nlm.nih.gov/BLAST/blast_FAQs.html. One of skill in the art will appreciate that these sequence identity ranges are provided for guidance only; it is entirely possible that strongly significant homologs could be obtained that fall outside of the ranges provided.

Recombinant: A recombinant nucleic acid is one that has a sequence that is not naturally occurring or has a sequence that is made by an artificial combination of two otherwise separated segments of sequence. This artificial combination is often accomplished by chemical synthesis or, more commonly, by the artificial manipulation of isolated segments of nucleic acids, e.g., by genetic engineering techniques.

cDNA (complementary DNA): A piece of DNA lacking internal, non-coding segments (introns) and regulatory sequences that determine transcription. cDNA is synthesized in the laboratory by reverse transcription from messenger RNA extracted from cells.

ORF (open reading frame): A series of nucleotide triplets (codons) coding for amino acids without any termination codons. These sequences are usually translatable into a peptide.

Operably linked: A first nucleic acid sequence is operably linked with a second nucleic acid sequence when the first nucleic acid sequence is placed in a functional relationship with the second nucleic acid sequence. For instance, a promoter is operably linked to a coding sequence if the promoter affects the transcription or expression of the coding sequence. Generally, operably linked DNA sequences are contiguous and, where necessary to join two protein-coding regions, in the same reading frame.

Subject: Living multi-cellular vertebrate organisms, a category that includes both human and non-human mammals.

Mimetic: A molecule (such as an organic chemical compound) that mimics the activity of a protein, such as calreticulin and therapeutically effective variants and fragments thereof. Peptidomimetic and organomimetic embodiments are within the scope of this term, whereby the three-dimensional arrangement of the chemical constituents of such peptido- and organomimetics mimic the three-dimensional arrangement of the peptide backbone and component amino acid sidechains in the peptide, resulting in such peptido- and organomimetics of the peptides having substantial specific inhibitory activity. For computer modeling applications, a pharmacophore is an idealized, three-dimensional definition of the structural requirements for biological activity. Peptido- and organomimetics can be designed to fit each pharmacophore with current computer modeling software (using computer assisted drug design or CADD). See Walters, "Computer-Assisted Modeling of Drugs", in Klegerman & Groves, eds., 1993, *Pharmaceutical Biotechnology*, Interpharm Press: Buffalo Grove, IL, pp. 165-174 and *Principles of Pharmacology* (ed. Munson, 1995), chapter 102 for a description of techniques used in computer assisted drug design.

As noted above, the present invention is based on the discovery that calreticulin and certain fragments of this protein, including the $\Delta 120$ calreticulin (Seq. I.D. No. 9), the 180 amino acid N-terminal domain ("vasostatin"; Seq. I.D. No. 4), and the 49 (Seq. I.D. No. 6), 60 (Seq. I.D. No. 8), and 61 (Seq. I.D. No. 5) amino acid fragments have one or more of the following biological

activities: (1) the ability to inhibit endothelial cell growth; (2) the ability to inhibit angiogenesis; and (3) the ability to suppress tumor growth. The following discussion explains how these activities were discovered and characterized.

5 **II. Calreticulin and Fragments of Calreticulin**
 Inhibit Endothelial Cell Proliferation

A. Endothelial Cell Assays

 Human and bovine endothelial cells were used for growth inhibition assays, and were
10 prepared as follows:

 Fetal bovine heart endothelial cells (American Type Culture Collection, ATCC, Manassas, VA) were grown through passage 12 in DMEM culture medium (BioWhittaker, Walkersville, MD) containing 10% heat inactivated fetal bovine serum (BioWhittaker), 100 ng/ml bFGF (R&D Systems, Minneapolis, MN), and 5 µg/ml gentamicin (Sigma). For proliferation assays, cells were trypsinized
15 (Trypsin/EDTA, Gibco BRL), washed, suspended in culture medium (DMEM containing 10% heat inactivated fetal bovine serum and 5 µg/ml gentamicin), plated (800 cells/well in 0.2 ml culture medium) in triplicate onto 96 well plates, and incubated for 5 days. DNA synthesis was measured by ³H thymidine deoxyribose uptake (0.5 mCi/well, 6.7 Ci/mmol; New England Nuclear, Boston, MA) during the last 20-23 hours of culture; cells were detached from the wells by freezing and thawing
20 (Angiolillo et al., *J. Exp. Med.* 182:155-162, 1995)

 Human umbilical vein endothelial cells were prepared from umbilical cord by 0.1% collagenase II (Worthington Biochemical Co., Freehold, NJ) digestion, as described (Gordon et al., *In Vitro* 19:661-671, 1983), and were grown through passage 5 in M199 culture medium (Sigma) supplemented with 20% newborn calf serum (Sigma), 5% human AB serum, 1.6 mM L-glutamine
25 (GibcoBRL), 50 µg/ml porcine heparin (Sigma), 50 µg/ml ascorbate (Fisher, Fairlawn, NJ), 15 mM HEPES buffer (Calbiochem-Behring, La Jolla, CA), and 15 µg/ml Endothelial Cell Growth Supplement (ECGF; a crude extract of bovine neural tissue containing basic and acidic FGF, Sigma). Endothelial cell purity was greater than 95%, as determined by staining with a rabbit antiserum to human Factor VIII-related antigen (Dako, Carpinteria, CA). For proliferation assays, cells were
30 trypsinized, washed, suspended in culture medium (RPMI 1640, BioWhittaker) supplemented with 18% heat-inactivated fetal bovine serum (BioWhittaker) and 18 units/ml porcine heparin (Sigma), and plated (3.5×10^3 cells/well in 0.2 ml culture medium) in triplicate cultures into 96-well plates. After incubation for 72 hours, DNA synthesis was measured by ³H thymidine deoxyribose uptake during the last 20-23 hours of culture.

35

**B. Identification of Calreticulin
as an Endothelial Cell Growth Inhibitor**

Using inhibition of bFGF (basic Fibroblast Growth Factor) -induced endothelial cell proliferation as an assay to monitor inhibitory activity, inhibitory compounds were purified from serum-free culture supernatants of the EBV-immortalized cell line VDS-O (maintained as described by Tosato, et al., *J. Immunol.* 137:2037-2042, 1986). For production of conditioned medium, exponentially growing VDS-O cells were washed free of serum, and cultured for 48 hours at the concentration of 2.0×10^6 cells/ml in serum free PFHM-11 medium (GIBCO/BRL, Grand Island, NY) supplemented with 5 µg/ml gentamicin (Sigma Chemical Co., St. Louis, MO). At the end of incubation, cells and debris were removed by centrifugation and sterile filtration (0.45 µ filters), and 6.0 µg/ml Aprotinin (Sigma) was added to the conditioned medium.

After clarification, filtration and concentration, the culture supernatant was purified by four sequential chromatographic steps, including metal chelating affinity, anion exchange, hydrophobic interaction, and anion exchange. Silica gel 60 (EM Science, Gibbstown, New Jersey), used for absorption of non-polar substances, was added to the conditioned medium at the concentration of 5 gm/l. After rotation, the silica was removed, and the conditioned medium was filtered (Ultrapump II, Filtron Technology Corp, Northborough, MA) through a membrane with an 8,000 dalton MW cutoff (Ultrasette Omega, Filtron). Concentrated (15 fold) supernatant was exchanged into 10 mM $\text{Na}_2\text{HPO}_4/\text{NaH}_2\text{PO}_4$, 100 mM NaCl, 0.1mM imidazole (Sigma) at pH 8.3, and applied to a Chelating Sepharose Fast Flow column (2.5 x 30 cm; Amersham Pharmacia Biotech, Piscataway, NJ) equilibrated in the same buffer. Bound material was eluted with a 10 mM $\text{Na}_2\text{HPO}_4/\text{NaH}_2\text{PO}_4$, 100 mM NaCl buffer containing 50 mM imidazole. Active fractions were equilibrated into 20 mM $\text{Na}_2\text{HPO}_4/\text{NaH}_2\text{PO}_4$ buffer (pH 7.8), and applied to an anion exchange Resource Q column (6 ml; Amersham Pharmacia Biotech) equilibrated with the same buffer. Bound fractions were eluted with a linear gradient of 200 mM to 1.0 M NaCl in $\text{Na}_2\text{HPO}_4/\text{NaH}_2\text{PO}_4$ buffer, pH 6.25. Active fractions were adjusted to 1.2 M $(\text{NH}_4)_2\text{SO}_4$, and applied to Macro-Prep Methyl Hydrophobic Interaction Chromatography (HIC) column (2.5x30 cm; Bio-Rad Laboratories, Hercules, CA) equilibrated with 20 mM $\text{Na}_2\text{HPO}_4/\text{NaH}_2\text{PO}_4$ buffer containing 1.2 M $(\text{NH}_4)_2\text{SO}_4$, at pH 6.5. Bound material was eluted by a linear decreasing gradient of 1.2 to 0 M $(\text{NH}_4)_2\text{SO}_4$ in 20 mM $\text{Na}_2\text{HPO}_4/\text{NaH}_2\text{PO}_4$ buffer. Biologically active fractions were pooled and loaded onto a Mini Q PC 3.2/3 anion exchange column (Amersham Pharmacia Biotech) equilibrated with 20 mM $\text{Na}_2\text{HPO}_4/\text{NaH}_2\text{PO}_4$ buffer at pH 7.8, mounted on a Smart System (Amersham Pharmacia Biotech) equipped with a superloop. Bound material was eluted by a linear gradient of NaCl (20 mM to 1.0 M) in the starting buffer.

In a representative purification, an initial loading with 340 mg total protein from 16 liters of supernatant yielded 4.5 µg of purified protein. The total biologic activity of the starting supernatant was approximately 44×10^3 units (a unit being defined as the 1/2 maximal activity measured in a proliferation assay of fetal bovine heart endothelial cells), and the purified material contained

approximately 9×10^3 units, a recovery of approximately 26% of the original biologic activity. The specific activity (units/mg protein) of the purified material was estimated to be 2×10^6 units.

The biologically active material was analyzed by two-dimensional gel electrophoresis under reduced conditions followed by Coomassie staining (not shown). Two-dimensional polyacrylamide gel electrophoresis was performed as described (Wirth et al., *Electrophoresis* 16:1946-1960, 1995). Samples for electrophoresis were prepared in buffer containing 8 M urea, 20 mM Tris, pH 6.8, 30 mM DTT, 4% CHAPS, and 2% Pharmalytes, pH 3-10. Precast immobilized pH gradient gel strips formulated with nonlinear pH gradients, pH 3-10 NL (180 x 3 mm x 0.5 mm) were purchased from Pharmacia LKB and rehydrated overnight at room temperature in 8M urea, 10 mM DTT, 2% w/v CHAPS, and 2% Pharmalytes, pH 3-10. Second dimension SDS-PAGE was performed using 1.5 mm thick 10% SDS polyacrylamide gels at constant current (40 mA/gel) at 10 °C. For SDS-PAGE and Western blot analysis, protein was solubilized in Tricine SDS sample buffer (Novex San Diego, CA), boiled, and run through 10-20% Tricine Gels. Prestained molecular weight protein standards (range 4-250 kDa, Novex) were used throughout. Gels were stained with Colloidal Coomassie G-250 stain (Colloidal Coomassie kit, Novex) or silver stain (SilverXpress staining kit, Novex). Two well defined polypeptide spots were identified with approximate molecular weight of 55 and 20 kDa and apparent pI of 4.7 and 5.6, respectively. In addition, a series of poorly defined spots were visualized with relative molecular weights ranging between 30 and 40 kDa.

The well defined spots were subjected to trypsin digestion and the tryptic fragments were analyzed by ion trap mass spectrometry. By this method, the 55 kDa polypeptide was identified as human calreticulin and the 20 kDa polypeptide as the light chain of human ferritin.

Further evidence of the identity of the 55 kDa component to calreticulin was obtained by Western blotting of the purified material. After electrophoresis, protein was transferred onto Immobilon-P membranes (Millipore, Bedford, MA). For detection of calreticulin, the membranes were incubated overnight with a rabbit anti-human calreticulin antiserum (Affinity Bioreagents Inc., Golden, CO), a rabbit anti-calreticulin N, or anti-calreticulin P domain antiserum (Pogue et al., *J. Virol.* 67:7106-7117, 1993). Bound antibody was detected with an affinity-purified, peroxidase-linked, donkey anti-rabbit IgG antibody (Amersham Pharmacia Biotech, Inc., Piscataway, NJ) and a chemiluminescence detection system (ECL kit, Amersham Pharmacia Biotech).

To gain further insight on the nature of these calreticulin fragments, antisera for human calreticulin N-terminal (aa 6-19) and C-terminal (aa 382-400) peptides were used in immunoblotting (Pogue et al., *J. Virol.* 67:7106-7117, 1993). Both these antisera identified the 55 kDa band and thus confirmed its identity to calreticulin. However, only the antiserum to the N-terminal calreticulin peptide reacted with the 30-40 kDa bands. Therefore, the biologically active, purified material from culture supernatant of the VDS-O cell line contained human calreticulin, the light chain of human ferritin, and N-terminal fragments of calreticulin. This result led to the investigation of other activities of calreticulin and calreticulin fragments, including the 180 amino acid N-terminal domain

fragment ("vasostatin"; Seq. I.D. No. 4), Δ 120 calreticulin (Seq. I.D. No. 9), the 49 (Seq. I.D. No. 6), 60 (Seq. I.D. No. 8) and 61 (Seq. I.D. No. 5) amino acid fragments, as described below.

III. Production of Recombinant Calreticulin and Calreticulin Fragments

5 The expression of human calreticulin, the calreticulin N-terminal deletion fragment lacking amino acids 1-120 (Seq. I.D. No. 9), and the N-terminal domain (Seq. I.D. No. 4) fused to maltose-binding protein (MBP) in *Escherichia coli* was previously reported (Singh et al., *Proc. Natl. Acad. Sci. U.S.A.* 91:12770-12774, 1994; Atreya et al., *J. Virol.* 69:3848-3851, 1995). *E. coli* cells containing the clones described in Singh et al., *Proc. Natl. Acad. Sci. U.S.A.* 91:12770-12774, 1994 and Atreya et al., *J. Virol.* 69:3848-3851, 1995 were grown in Luria-Bertani broth (Advanced Biotechnologies, Inc., Columbia MD) with 0.2% glucose and 100 μ g/ml ampicillin (Sigma) to an OD₆₀₀ of approximately 1.0, and fusion protein expression was induced with 0.3 mM IPTG (GIBCO/BRL) for 2-2.5 hours. After lysis (1 μ g/ml lysozyme in 10 mM Tris, pH 7.5, 5% glycerol, 100 mM EDTA, 5 mM B-ME), sonication, and centrifugation (30 min at 8360 x g) of the bacterial suspension, the supernatant was loaded onto a preequilibrated (20 mM Tris, pH 7.5, 200 mM NaCl, and 1 mM EDTA) 15 ml amylose column (New England Biolabs, Beverly, MA), and the bound material was eluted with 10 mM Maltose. Protein-containing fractions were ultracentrifuged (2 hours at 104,000 x g); supernatant was retained. Separation of MBP from calreticulin (Seq. I.D. No. 2) and vasostatin (Seq. I.D. No. 4) was accomplished by cleavage with factor Xa (New England Biolabs, Beverly, MA). Purification of cleaved calreticulin (Seq. I.D. No. 2) or vasostatin (Seq. I.D. No. 4) from MBP was achieved by anion exchange chromatography using a preequilibrated (20 mM Tris, pH 8.0, 25 mM NaCl) Resource Q column (Amersham Pharmacia Biotech). Bound material was eluted by a step-wise gradient where MBP elutes at 100-150 mM NaCl; Factor Xa elutes at approximately 400 mM NaCl; and calreticulin or vasostatin elute at approximately 250 mM NaCl.

25 For expression of the calreticulin fragment encompassing amino acids 120-180 (Seq. I.D. No. 5), the coding region for this fragment was amplified by PCR and then cloned (confirmed by sequencing) as an N-terminal fusion protein with the MBP gene for expression in *E. coli*. The growth of *E. coli* and protein purification were performed as described above for MBP-calreticulin.

A recombinant form of calreticulin fused to glutathione S-transferase (GST) was also produced. For construction of the GST-calreticulin fusion construct, the coding region for the mature calreticulin protein was cloned as a C-terminal translational fusion with the glutathione S transferase (GST) gene for expression in *E. coli*. The growth of *E. coli*, the induction and release of GST-calreticulin (Seq. I.D. No. 2) from the bacteria was the same as described above for MBP-calreticulin, except for IPTG induction (0.6 mM). For purification of GST-calreticulin, the bacteria were sonicated, pelleted, and the supernatants (adjusted to pH 7.0) were mixed with prewashed Glutathione Sepharose 4B (Bulk GST purification module; Amersham Pharmacia Biotech) in PBS with 1.0% Triton X-100. After 30 min incubation and washing the beads, bound protein was eluted with a 50 mM Tris-HCl buffer containing 10 mM glutathione, pH 8.0. Eluted material was ultracentrifuged (2 hours at 104,000 x g), and supernatant retained. All protein lots for *in vivo* and *in vitro* experiments

(GST-calreticulin, control GST, MBP-calreticulin, MBP-vasostatin, MBP, cleaved calreticulin and vasostatin) were tested for endotoxin by the Limulus Amebocyte Lysate (LAL) kinetic-QCLTM assay (BioWhittaker) and were found to contain <5 units/10 µg protein.

5 The purified recombinant proteins including the expression tag sequences were resolved by SGS-PAGE as discrete bands migrating at the expected relative positions. Calreticulin, cleaved from MBP-calreticulin by treatment with Factor Xa and subsequently purified by anion exchange chromatography, resolved as a doublet with a relative molecular weight of approximately 50 and 55 kDa.

10 Natural calreticulin was also obtained from purified B cell line supernatant by eluting the band at 55 kDa from a preparative Tris-glycine gel. SDS PAGE analysis and silver staining documented the isolation of a doublet at approximately 55 kDa from other copurified components.

When tested in functional assays, recombinant purified MBP-calreticulin (Seq. I.D. No. 2) inhibited the proliferation of fetal bovine heart endothelial cells induced by bFGF. At a concentration of 1 µg/ml, MBP-calreticulin (Seq. I.D. No. 2) inhibited fetal bovine heart endothelial cell growth by 67% while control MBP had minimal effects (Figure 1). Similar inhibition was noted with 1 µg/ml GST-calreticulin (Table 1). In addition, recombinant calreticulin (Seq. I.D. No. 2) that had been cleaved and purified from MBP-calreticulin, as well as purified natural calreticulin inhibited the proliferation of fetal bovine heart endothelial cells, while control MBP did not (Table 1). By contrast, recombinant purified light chain of human ferritin (a gift from Dr. P. Arosio, San Raffaele, Milan, Italy) had minimal effect on the proliferation of endothelial cells when tested at 1 pg to 1 µg/ml (not shown).

To assess whether specific fragments of calreticulin exhibit inhibitory activity, the components from cell line culture supernatants with a relative molecular weight of 30-40 kDa which were immunoreactive with antisera to calreticulin were isolated by gel elution. A component with a relative molecular weight of approximately 34 kDa was separated from calreticulin at 55-60 kDa. This gel-eluted calreticulin fragment inhibited endothelial cell proliferation (Table 1). Thereafter, the N-terminal calreticulin domain (Seq. I.D. No. 4), including amino acids 1-180 of the mature protein, was produced in *E. coli* as a fusion protein of MBP (MBP-calreticulin-N, Atreya et al., *J. Virology* 69:3848-3851, 1995). The purified recombinant MBP-calreticulin N-terminal domain and the cleaved calreticulin N-terminal domain inhibited the proliferation of fetal bovine heart and of human umbilical vein endothelial cells (Table 2).

Table 1. Inhibition of Endothelial Cell Proliferation by Calreticulin

	Additions to Culture	Proliferation mean cpm/culture (\pm SD)	Inhibition (%)
5	1. None	5,731 (2344) ^a	
	bFGF	23,243 (8630)	
	GST-Calreticulin	12,503 (4510)	46
	GST	25,410 (8158)	
10	2. None	5,510 (2040) ^b	
	bFGF	25,322 (7849)	
	Cleaved calreticulin	7,088 (1846)	72
	Cleaved MBP	21,410 (232)	
15	3. None	15,277 ^c	
	bFGF	57,613	
	Gel-eluted calreticulin	11,226	81
	Gel-eluted calreticulin fragment	32,039	45
	Gel-eluted control	67,796	

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Fetal bovine heart endothelial cells (800 cells/well) were cultured for 5 days in medium alone or medium supplemented with bFGF (25 ng/ml). Recombinant purified GST-calreticulin (Seq. I.D. No. 2; 1 μ g/ml), control recombinant GST (1 μ g/ml), calreticulin cleaved and purified from MBP-calreticulin (Seq. I.D. No. 3; 1 μ g/ml), MBP cleaved and purified from MBP-calreticulin (1 μ g/ml), natural calreticulin (estimated concentration 0.160 μ g/ml) or natural fragments of calreticulin (estimated concentration 0.8 μ g/ml) derived by gel elution of calreticulin (purified from cell line culture supernatant) were added to endothelial cell cultures with bFGF (25 ng/ml). Proliferation was measured by ³H thymidine incorporation during the final 20-23 hours of culture; the results reflect mean cpm/culture.

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30 ^a Reflects the mean of 9 experiments, each performed in triplicate

^b Reflects the mean of 2 experiments, each performed in triplicate

^c Reflects the mean of triplicate cultures; SDs within 12% of the mean

Table 2. Inhibition of Endothelial Cell Proliferation by Vasostatin

	Additions to Culture	Proliferation (mean cpm/culture)	Inhibition (%)
35	1. ^a None	142	
	bFGF	32,493	
	MBP-vasostatin	17,930	44.8
40	MBP	34,363	
	Vasostatin ^c	13,231	59.3
	2. ^b None	4,210	
	bFGF	28,050	
45	Vasostatin	9,864	64.8

Human umbilical vein endothelial cells (3.5×10^3 cells/well) or fetal bovine heart endothelial cells (800 cells/well) were cultured in medium alone or in medium supplemented with bFGF. Recombinant purified vasostatin (1 μ g/ml), MBP-vasostatin (1 μ g/ml), or MBP (1 μ g/ml) were added

to bFGF-supplemented cultures. Proliferation was measured by ^3H thymidine incorporation during the final 20-23 hours of culture.

^a Cultures containing human umbilical cord derived endothelial cells; bFGF used at 25 ng/ml. The results reflect the mean of triplicate cultures; SDs within 10% of the mean.

5 ^b Cultures containing fetal bovine heart endothelial cells; bFGF 25 ng/ml. The results reflect the mean of 3 experiments; SDs within 12% of the mean.

^c Calreticulin is identified as Seq. I.D. No. 4.

10 Smaller fragments of the calreticulin N-terminal domain were then synthesized. These fragments were tested for activity against fetal bovine heart endothelial cells as described above. Both were found to be active, with the 49 amino acid fragment (Seq. I.D. No. 6) producing 45-69% inhibition at a concentration of 50 $\mu\text{g/ml}$ and the 60 amino acid fragment (Seq. I.D. No. 8) producing 59.5-88% inhibition at a concentration of 100 $\mu\text{g/ml}$ (Table 3). It should be noted that a control
15 peptide 60 amino acids in length (from the chemokine RANTES) which was similarly synthesized and purified, had no inhibitory effect in these assays. It should also be noted that the levels of endothelial cell growth inhibition by the 49 (Seq. I.D. No. 6) and the 60 (Seq. I.D. No. 8) amino acid fragments were comparable to those derived from vasostatin (Seq. I.D. No. 4). The higher amounts of peptide being required to achieve endothelial cell growth inhibition compared to vasostatin (Seq.
20 I.D. No. 4) are likely attributable to the differences in the manufacturing process resulting in peptide denaturing and/or unfavorable folding. To ensure that growth inhibition of endothelial cell growth *in vitro* was not the result of non-specific toxicity, the 49 (Seq. I.D. No. 6) and the 60 (Seq. I.D. No. 8) amino acid fragments were tested for their effects on the proliferation of 2 indicator B cells lines, VDS-O and CA-46. At concentrations ranging between 1-100 $\mu\text{g/ml}$, the 49 (Seq. I.D. No. 6) and the
25 60 (Seq. I.D. No. 8) amino acid fragments had minimal effect on the proliferation of these cell lines.

To further ensure that fragments of calreticulin are active as inhibitors of endothelial cell growth, recombinant MBP- Δ 120 calreticulin (calreticulin missing the N-terminal 1-120 amino acids; Seq. I.D. No. 9) and MBP-120-180 calreticulin (Seq. I.D. No. 5) were tested. Recombinant purified MBP- Δ 120 calreticulin (Seq. I.D. No. 9) inhibited the proliferation of fetal bovine heart endothelial
30 cells, and the degree of inhibition was comparable to that of calreticulin (Figure 4).

Vasostatin (calreticulin N-domain, encompassing amino acids 1-180; Seq. I.D. No. 4), full length calreticulin (Seq. I.D. No. 2), calreticulin minus 17 amino acids from the N-terminus (Seq. I.D. No. 3), and Δ 120 calreticulin (Seq. I.D. No. 9) inhibited bFGF-induced endothelial cell proliferation to a similar degree at similar concentrations. Hence, the activity of the 61 amino acid calreticulin
35 fragment (Seq. I.D. No. 5) was also determined. This was done by purifying the MBP-120-180 calreticulin fragment (Seq. I.D. No. 5) from *E. coli* expressing the recombinant protein fused to MBP (Figure 5A). When tested for inhibition of endothelial cell proliferation, recombinant purified MBP-120-180 calreticulin fragment (Seq. I.D. No. 5) inhibited the proliferation of bovine heart endothelial cells induced by bFGF (Figure 5B). At a concentration of 0.5 $\mu\text{g/ml}$, the 61 amino acid calreticulin
40 fragment (Seq. I.D. No. 5) inhibited fetal bovine heart endothelial cell proliferation by 65%. A side-

by-side comparison of full length MBP-calreticulin (Seq. I.D. No. 3), MBP-vasostatin (Seq. I.D. No. 4), MBP- Δ 120 calreticulin (Seq. I.D. No. 9) and MBP-120-180 calreticulin (Seq. I.D. No. 5) fragment revealed that, on a molar basis, the four proteins display similar endothelial cell growth inhibitory active *in vitro*. Control MBP was not inhibitory (Figure 6).

- 5 Hence, both the calreticulin fragment identified as Seq. I.D. No 5 (the 61 amino acid peptide) and the calreticulin fragment identified as Seq. I.D. No 6 (the 49 amino acid peptide) specifically inhibit endothelial cell growth *in vitro*.

Table 3 Inhibition of Endothelial Cell Proliferation by Calreticulin 60 and 49 amino acid fragments

	Addition to Culture	Proliferation (mean cpm/culture)	Inhibition (%)
5	1. None	5,290 ^a	
	bFGF	13,950	
	MBP-vasostatin	8,585	38.5
	60 amino acid peptide (100 µg/ml) ^b	1,688	87.9
10	60 amino acid peptide (50 µg/ml)	8,318	40.1
	49 amino acid peptide (50 µg/ml) ^c	4,368	68.7
	49 amino acid peptide (25 µg/ml)	6,037	56.7
	49 amino acid peptide (12.5 µg/ml)	9,743	30.2
	RANTES (100 µg/ml)	16,443	
15	2. None	4,434	
	bFGF	9,160	
	60 amino acid peptide (100 µg/ml)	3,709	59.5
	49 amino acid peptide (50 µg/ml)	5,004	45.4
20	RANTES (50 µg/ml)	10,291	
	RANTES (12.5 µg/ml)	13,644	

Fetal bovine heart endothelial cells (800 cells/well) were cultured for 5 days in medium alone or medium supplemented with bFGF (25 ng/ml). Recombinant MBP vasostatin (0.5 µg/ml), a synthetic calreticulin 60 amino acid peptide (50 and 100 µg/ml), a synthetic calreticulin 49 amino acid peptide (12.5-50 µg/ml), and a synthetic 60 amino acid peptide of the chemokine RANTES (12.5-100 µg/ml) were added to endothelial cell cultures with bFGF (25 µg/ml). Proliferation was measured by ³H thymidine incorporation during the final 20-23 hours of culture; the results reflect the mean cpm/culture.

^a Reflects the mean of triplicate cultures; SDs within 10% of the mean.

^b The 60 amino acid calreticulin peptide (amino acids 121-180) is identified as Seq. I.D. No 8

^c The 49 amino acid calreticulin peptide (amino acids 132-180) is identified as Seq. I.D. No. 6

IV. Calreticulin and Calreticulin Fragments Do Not Inhibit Growth of Nonendothelial Cells

Following the observation that calreticulin and calreticulin fragments inhibit the proliferation of endothelial cells, these proteins were then assayed for inhibitory effect on non-endothelial cell types.

Human mononuclear cells, and mononuclear cells enriched for T or B lymphocytes, were used in proliferation assays in the presence of phytohemagglutinin (Sigma), pokeweed-mitogen (Sigma), or Epstein-Barr virus (B95-8 strain), as described (Tosato et al., *J. Immunol.* 140:4329-4336, 1988). The lymphoblastoid cell line VDS-O (Tosato, et al., *J. Immunol.* 137:2037-2042, 1986); the Burkitt lymphoma cell lines CA46, BL-41, KK124, Ag876, SHO (Cherney et al., *Cancer Res.*

57:2508-2515, 1997); the T-cell line Molt-4 (ATCC); the Hodgkin's lymphoma cell line Hs445 (ATCC); the prostate adenocarcinoma cell lines TSU-Pr1 (from A. Passaniti, NIH, Baltimore, MD), Du145 (ATCC), and PC3 (ATCC); and the acute promyelocytic leukemia cell line HL-60 (ATCC) were cultured in RPMI 1640 medium with 10% heat inactivated fetal calf serum (BioWhittaker), 20 mM L-glutamine (Sigma) and 5 µg/ml gentamicin, and tested in 3-5 day proliferation assays at cell densities ranging from 300 to 2400 cells/ well. All other cell lines were tested for proliferation in 3-day assays. The neuroblastoma cell line SKNMC (ATCC) was cultured in EMEM medium (BioWhittaker) with 10% heat inactivated fetal calf serum (BioWhittaker) and 5 µg/ml gentamicin (Sigma) and tested for proliferation at 125-1000 cells/well. The lung adenocarcinoma cell line A549 (ATCC) was cultured in F-12 Nutrient Mixture (HAM, GIBCO BRL) with 10% heat inactivated fetal calf serum (BioWhittaker) and 5 µg/ml gentamicin (Sigma), and tested for proliferation at 250-2000 cells/well. The breast adenocarcinoma MDA-MB-468 (ATCC) and the Wilms tumor SK-NEP-1(ATCC) cell lines were cultured in Leibovitz L-15 medium (GIBCO BRL) supplemented with 10% heat inactivated fetal calf serum (BioWhittaker) and 5 µg/ml gentamicin (Sigma), and tested for proliferation at 300-2500 cells/well. The colon carcinoma cell line SW480 (ATCC), the melanoma cell line A-375 (ATCC), and human foreskin fibroblasts (H5 68, ATCC) were cultured in DMEM medium (BioWhittaker) with 10% heat inactivated fetal calf serum (BioWhittaker) and 5 µg/ml gentamicin (Sigma), and were tested for proliferation at 500-4000 cells/well. All cell lines tested mycoplasma negative.

The results of these assays showed that, in contrast to their inhibitory effect on human and bovine endothelial cell growth, MBP-calreticulin (Seq. I.D. No. 3) and the MBP-calreticulin N-terminal 180 amino acid fragment (Seq. I.D. No. 4) had minimal effect on the proliferation of a variety of other primary cells and cell lines, when administered at concentrations of 0.5-2.5 µg/ml.

The 61 amino acid fragment of calreticulin (MBP-calreticulin 120-180; Seq. I.D. No. 5), as well as the 60 and 49 amino acid peptides of calreticulin (Seq. I.D. No 8, and 6, respectively) were also tested for their ability to inhibit the proliferation *in vitro* of cells other than endothelial cells. At concentrations ranging between 1 and 100 µg/ml, neither peptide inhibited the proliferation of the human B cell lines VDS-O and CA-46. Thus, calreticulin (Seq. I.D. No. 3), the calreticulin N-terminal 180 amino acid fragment, vasostatin (Seq. I.D. No. 4), a 60 amino acid calreticulin peptide included in the N-terminal amino acid peptide (amino acids 121-180; Seq. I.D. No. 8), and a 49 amino acid calreticulin peptide (amino acids 132-180; Seq. I.D. No. 6) included in the N-terminal 180 amino acid peptide specifically inhibit endothelial cell proliferation.

V. Calreticulin and Calreticulin Fragments Inhibit Angiogenesis

The murine Matrigel assay (Passaniti et al., *Lab. Invest.* 67:519-528, 1992) was employed to evaluate the effects of calreticulin and calreticulin fragments on angiogenesis *in vivo*. The Matrigel assay was performed as described by Angiolillo et al. (*J. Exp. Med.* 182:155-162, 1995). Matrigel, a crude extract of the Englebreth-Holm-Swarm tumor, was obtained from Collaborative Biomedical Products, Becton Dickinson Labware, Bedford, MA. An aliquot (0.5 ml) of Matrigel alone or with

desired additives was injected subcutaneously (s.c.) into the mid abdominal region of female BALB/c nude mice, 6-8 weeks old. Five mice were injected with each mixture. After 5-7 days, the animals were sacrificed, Matrigel plugs were removed, fixed in 10% neutral buffered formalin solution (Sigma), and embedded in paraffin. Tissues were sectioned (5 μ thickness), and slides stained with

5 Masson's trichrome. Quantitative analysis of angiogenesis in Matrigel plugs utilized a computerized semi-automated digital analyzer (Optomax, 40-10 System, Hollis, NH). The instrument was adjusted to evaluate a circular area measuring 1.26×10^5 mm² of Matrigel, and within this area, to measure the area occupied by cells. For each plug, 12-15 distinct fields were evaluated. The fields were

10 randomly selected from each plug, and the operator was blind to the experimental design. The average area occupied by cells/ 1.26×10^5 mm² Matrigel field was calculated. Results are expressed as the mean area occupied by cells/Matrigel field.

Matrigel plugs were fixed in 10% neutral buffered formalin solution (Sigma), embedded in paraffin, sectioned at 4 μ m, and stained with Masson's trichrome by standard methods.

When added to Matrigel at concentrations of 1.25-5 μ g/ml, GST-calreticulin (Seq. I.D. No. 2) displayed a concentration-dependent inhibition of bFGF-induced neovascularization (Table 4, exp. 1). Similarly, at 5-10 μ g/ml, the MBP-1-180 amino acid fragment (termed "vasostatin" because of this activity; Seq. I.D. No. 4) inhibited bFGF-induced neovascularization (Table 4, exp. 2). In

15 addition, at 5 μ g/ml, vasostatin (cleaved and purified from MBP-vasostatin) inhibited bFGF-induced Matrigel vascularization comparably to MBP-vasostatin and MBP-calreticulin (Table 4, exp. 3).

20 Furthermore, MBP- Δ 120 calreticulin (Seq. I.D. No. 9) and 120-180 calreticulin (Seq. I.D. No. 5) inhibited bFGF-induced Matrigel vascularization comparably to cleaved recombinant calreticulin (Table 4, exp. 4).

Table 4. Effects of Vasostatin and Calreticulin on Angiogenesis *in vivo*

Exp. #	Additions to Matrigel	Mean Surface Area Occupied by Cells (mm ² /1.26x10 ⁵ mm ²)	Inhibition (%)
5	1. None	671	
	bFGF	17,732	
	bFGF+GST-calreticulin (5 µg/ml)	4,616	74
	bFGF+GST-calreticulin (2.5 µg/ml)	6,387	64
	bFGF+GST-calreticulin (1.25 µg/ml)	9,870	44
10	2. None	649	
	bFGF	11,544	
	bFGF+MBP-vasostatin (10 µg/ml)	4,539	61
	bFGF+MBP-vasostatin (5 µg/ml)	5,286	54
15	bFGF+MBP (10 µg/ml)	9,186	20
	3. None	487	
	bFGF	14,472	
	bFGF+MBP-calreticulin (5 µg/ml)	5,112	65
20	bFGF+MBP-vasostatin (5 µg/ml)	4,989	66
	bFGF+vasostatin (5 µg/ml)	4,638	68
	MBP (5 µg/ml)	13,472	7
	4. None	1,329	
25	bFGF	8,320	
	bFGF+cleaved calreticulin	3,464	58
	bFGF+MBP-Δ120 calreticulin	4,291	48
	bFGF+MBP-120-180	2,458	70

30 Mice were injected subcutaneously with Matrigel alone, Matrigel plus bFGF (150 ng/ml), Matrigel plus bFGF (150 ng/ml) plus GST-calreticulin (Seq. I.D. No. 2), MBP-calreticulin (Seq. I.D. No. 3), MBP-vasostatin (Seq. I.D. No. 4), MBP, vasostatin, MBP-Δ120 calreticulin (Seq. I.D. No. 9), or MBP-120-180 calreticulin (Seq. I.D. No. 5). Plugs were removed after 5-7 days, and histologic sections were stained with Masson's trichrome. The results reflect the mean surface area (expressed in mm²) occupied by cells within a circular surface area of 1.26x10⁵ mm²; 12-15 non-overlapping fields were scanned in each plug; there were 5 plugs/group. Determinations of surface area were performed by a semi-automated digitalized analyzer.

40 VI. Calreticulin and Calreticulin Fragments Inhibit Tumor Growth

Calreticulin and calreticulin fragments were initially tested for their ability to prevent the growth of human Burkitt lymphomas in athymic mice.

BALB/c nu/nu female mice, 6 weeks of age (National Cancer Institute, Frederick, MD) maintained in pathogen-limited conditions, received 400 rad (1rad=0.01 Gy) total body irradiation and 24 hours later were injected s.c. in the right abdominal quadrant with 10⁷ exponentially growing human Burkitt lymphoma cells (CA46 cell line; Sgadari et al., *Proc. Natl. Acad. Sci. U.S.A.* 93:13791-13796, 1996) in 0.2 ml PBS. Immediately after the Burkitt cells were inoculated s.c., and continuing daily thereafter 6 days/week, the mice received s.c. injections proximal to the site of original cell inoculation of test calreticulin samples or appropriate controls. These included purified

GST-calreticulin (Seq. I.D. No. 2), control GST, MBP-calreticulin (Seq. I.D. No. 3), MBP-vasostatin (Seq. I.D. No. 4), MBP, or formulation buffer used to dilute test proteins (sterile saline solution containing 50 mg/ml human albumin and 5 mg/ml mannitol; endotoxin <5 units/ml). Tumor size was estimated (in cm²) twice weekly as the product of two-dimensional caliper measurements (longest perpendicular length and width). A subcutaneous mass appearing at or proximal to the site of cell inoculation was considered a tumor when it measured at least 0.16 cm² in surface area and increased in size by at least 0.1 cm² over the following week.

As expected, 12/12 mice injected with control GST protein developed a tumor by day 17. By contrast, only 4/13 mice injected with GST-calreticulin (Seq. I.D. No. 2) developed a tumor by day 17 (p=0.005)(Figure 3A). The tumor-bearing mice were sacrificed, and the remaining 9 non-tumor bearing mice were maintained untreated. Tumors eventually developed in 8 of the 9 mice that had received initial calreticulin treatment. The latest tumor developed on day 36, 22 days after treatment had ended. One mouse remains tumor free (>100 days).

Vasostatin (Seq. I.D. No. 4) was tested for its ability to prevent or delay Burkitt tumor growth in the same murine tumor model. In a representative experiment (3 performed), MBP-vasostatin (Seq. I.D. No. 4) was inoculated daily, 6 days/week, for 14 days at a dose of 30 µg/mouse (Figure 3B). At the end of treatment, 3/9 mice inoculated with MBP-vasostatin (Seq. I.D. No. 4) had developed a tumor, as opposed to 8/8 mice inoculated with control MBP (p=0.009). After treatment was suspended on day 14, all mice were maintained untreated. All but one mouse (tumor free for >100 days) developed a tumor; the latest tumor appeared 15 days after treatment had ended.

The effects of extended MBP-vasostatin (Seq. I.D. No. 4) treatment on tumor development were examined (Figure 3C). All animals received treatment with MBP-vasostatin (60 µg/mouse, daily, 6 days/week; (Seq. I.D. No. 4) for at least 18 days, at which time all animals with tumors were sacrificed. The remaining tumor-free animals continued treatment until tumor formation. By day 18, 4/12 animals treated with MBP-vasostatin (Seq. I.D. No. 4) as opposed to 12/12 control-treated animals had developed a tumor (p=0.0013). The mean (±SD) weight of tumors in the untreated control group (0.43 ±0.2 g) was greater than the weight of tumors from vasostatin-treated animals (0.21 ±0.05 g), but the weight difference did not reach statistical significance (p=0.059). With continued MBP-vasostatin (Seq. I.D. No. 4) treatment, one additional tumor appeared on day 23, but the remaining 7 animals remain tumor-free as of day 60.

In an additional experiment (Figure 3D), MBP-vasostatin (Seq. I.D. No. 4) was administered at a higher dose (100 µg/mouse, daily, 6 days/week) and compared it to a lower dose (20 µg/mouse, daily, 6 days/week). After 18 days of treatment, all mice (6/6) inoculated with formulation buffer alone had developed a tumor. By contrast, none of the mice (0/9) inoculated with MBP- vasostatin at the dose of 100 µg/mouse had developed a tumor (p=0.0002). In addition, 3/5 mice inoculated with MBP-vasostatin at the dose of 20 µg/mouse developed a tumor (not significantly different from control, p=0.018), indicating a dose effect. Treatment was continued unchanged until tumors appeared. As of day 30, only one tumor has appeared in the group treated with the highest dose.

Subsequent tests were done to determine the effect of MBP-vasostatin (Seq. I.D. No. 4) on established human Burkitt lymphoma. To this end, Burkitt tumors were first established in the animals and then treatment was initiated. As shown in Figure 4A, the rate of Burkitt lymphoma growth was significantly reduced in the group (9 mice) treated with MBP-vasostatin (Seq. I.D. No. 4) at a dose 200 µg/mouse compared to the controls (10 mice) treated with formulation buffer alone or MBP (p=0.003). Tumors were removed on day 48. The mean weight of Burkitt tumors in the control group (6.89±2.6 g) was significantly greater (p=0.0005) than the mean weight of tumors treated with MBP-vasostatin (2.74±1.6 g). In an additional experiment, MBP-vasostatin (Seq. I.D. No. 4) was used at a higher dose (200 µg/twice/day, 5 days/week and once/day 2 days/week) for 38 days. As shown in Figure 4B, MBP-vasostatin treatment promoted the disappearance of 4 out of 12 Burkitt tumors but, as expected, formulation buffer alone did not have this effect. After treatment was completed on day 38, all animals with tumor were sacrificed and tumor weight measured. The mean weight of Burkitt tumors in the control group (3.95±0.77 g) was significantly greater (p=0.014) than the mean weight of tumors treated with MBP-vasostatin (1.74±0.44 g; Seq. I.D. No. 4). All 4 mice treated with MBP-vasostatin (Seq. I.D. No. 4) and without a tumor on day 38 were observed without treatment. As of day 70, none of the tumors have reappeared. These experiments indicate that vasostatin can delay, prevent, or cure Burkitt tumor development in a murine model.

In an additional experiment, MBP-Δ120 calreticulin (30 µg/mouse; Seq. I.D. No. 9) was tested for its ability to prevent Burkitt tumor growth, and compared its effects to those of MBP-vasostatin (30 µm/mouse; Seq. I.D. No. 4). After 14 days of treatment, all mice (8/8) inoculated with control MBP (20 µg/mouse) developed a tumor. By contrast, only 3/9 mice inoculated with MBP-Δ120 calreticulin (Seq. I.D. No. 9), and 3/9 mice inoculated with MBP-vasostatin (Seq. I.D. No. 4) developed a tumor (p=0.009). After treatment was suspended on day 14, the mice were observed. One mouse from the group treated with MBP-Δ120 calreticulin (Seq. I.D. No. 9) and one mouse from the group treated with MBP-vasostatin (Seq. I.D. No. 4) remain tumor-free as of day 60. Thus, similar to MBP-vasostatin (Seq. I.D. No. 4), MBP-Δ120 calreticulin (Seq. I.D. No. 9) can prevent Burkitt tumor growth.

For histological examination, tumor plugs were fixed in 10% neutral buffered formalin solution (Sigma), embedded in paraffin, sectioned at 4 µm, and stained with hematoxylin and eosin, or Masson's trichrome by standard methods. Histology of tumors that emerged on MBP-calreticulin (Seq. I.D. No. 3) or MBP-vasostatin (Seq. I.D. No. 4) treatment showed subtle characteristic changes to the tumor vasculature, including intimal and medial thickening of the vessel wall. These alterations were absent from tumors of control animals. The tumor tissue from test and control animals was indistinguishable with respect to morphology of tumor cells and the number of mitoses. However, tumors from calreticulin and vasostatin -treated animals displayed occasional infiltration with neutrophils, histiocytes, and lymphocytes that were absent from control tumors. In addition, tumor tissues from animals treated with MBP-vasostatin or MBP-calreticulin displayed a significant reduction in the number of vessels identified by immunohistochemical staining for the endothelial

cell marker CD31. No abnormalities were noted on gross and histological examination of liver, spleen, kidneys, heart, lung and lymph nodes from MBP-calreticulin and MBP-vasostatin-treated animals.

5 **VII. Calreticulin Fragment Inhibits the Growth of Colon Carcinomas, Breast Adenocarcinomas and Neuroblastoma Tumors, Wilms Tumor, Rhabdomyosarcoma, Promyelomonocytic Lymphoma, Murine lymphoma, and Melanoma Tumors.**

Additional experiments were designed to test the effect of MBP-vasostatin treatment on human colon carcinoma established in athymic mice. To this end, BALB/c athymic mice were
10 inoculated s.c. with the cells of the human colon carcinoma SW-480 cell line. After a tumor appeared (at least 130 mm² in size), 12 mice were treated s.c. with MBP-vasostatin (Seq. I.D. No. 4) and 10 mice were similarly treated with formulation buffer. Tumor growth was significantly reduced in the group treated with MBP-vasostatin (Seq. I.D. No. 4) at a dose of 100 µg/mouse (12 mice) compared to the control group (10 mice) treated with formulation buffer alone (p=0.0003, Figure 8). All tumors
15 were removed on day 39 of treatment. The mean (±SD) weight of colon carcinoma tumors in the control group (3.04±0.6 g) was significantly (p=0.0004) greater than the weight of tumors from MBP-vasostatin-treated animals (1.48±0.64 g).

The effects of MBP-vasostatin on human neuroblastoma tumor growth in athymic mice were also tested. To this end, athymic BALB/c mice were inoculated s.c. with the human neuroblastoma
20 SK-N-MC cell line. Beginning on the day of cell inoculation and continuing thereafter daily, the mice received s.c. either control MBP (20 µg/day) or MBP-vasostatin (30 µg/day; Seq. I.D. No. 4). On day 30, 8 of the 12 mice receiving control MBP (mean tumor weight 1.74 g) and 5 of the 12 mice receiving MBP-vasostatin (mean tumor weight 1.78 g; Seq. I.D. No. 4) were sacrificed. Treatment was continued unchanged in the remaining animals until day 55 at which time all animals with tumor
25 were sacrificed. One of the 12 mice treated with MBP-vasostatin (Seq. I.D. No. 4) never developed a tumor and remains tumor-free without treatment as of day 220. As shown (Figures 9A and 9B), the rate of tumor growth was reduced in the mice treated with MBP-vasostatin (p=0.07). In addition, the mean (±SD) weight of neuroblastoma tumors removed on day 55 in the control group (8.579±1.21) was greater than the weight of tumors from MBP-vasostatin-treated animals (4.58±0.64 g).

Furthermore, the effect of MBP-vasostatin (Seq. I.D. No. 4) on human breast
30 adenocarcinoma growth in athymic mice was also tested (Figure 10). Athymic BALB/c nude mice were inoculated s.c. with the human breast adenocarcinoma cell line MDA-MB231. Beginning on the day of cell inoculation and continuing daily thereafter (5 days/week for 18 days) the mice were inoculated s.c. with either MBP-vasostatin (200 µg/day, 5 days/week and 100 µg/day 2 days week;
35 Seq. I.D. No. 4) or formulation buffer control. As shown, breast adenocarcinoma tumor growth was reduced by MBP-vasostatin. There was no evidence of local or systemic toxicity in vasostatin-treated animals.

To establish whether MBP-vasostatin could inhibit tumor growth in a syngeneic system, BALB/c nude mice were injected s.c. with the murine B-cell lymphoma cell line Raw 8.1 (also

BALB/c). Beginning on the day of cell inoculation and continuing thereafter daily, 9 mice were inoculated s.c. with control formulation buffer and 9 mice were treated with MBP-vasostatin (200 µg/day; Seq. I.D. No. 4). Treatment was stopped when all animals developed a tumor on day 15. As shown (Figure 11A), MBP-vasostatin treatment reduced significantly murine lymphoma tumor growth ($p < 0.01$). In addition (Figure 11B), the mean (\pm SD) weight of the tumors in the control group (4.51 ± 0.8 g) was significantly ($p < 0.01$) greater than the weight of tumors from MBP-vasostatin-treated animals (0.86 ± 0.34 g). These experiments demonstrate that MBP-vasostatin (Seq. I.D. No. 4) exhibits a potent antitumor effect not only in athymic mice bearing xenogeneic tumors, but also in athymic mice bearing syngeneic tumors.

Additional tests measuring the effects of MBP-vasostatin (Seq. I.D. No. 4) on Wilms tumors were done (Figure 12). Athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (12×10^6 cell/mouse) with the human Wilms tumor cell line (SK-NP-1). One day after cell injection, and continuing daily thereafter for 42 days, the mice received either control formulation buffer alone or MBP-vasostatin (100 µg/day, 6 days/week; Seq. I.D. No. 4). As shown, MBP-vasostatin (Seq. I.D. No. 4) induced a significant reduction of Wilms tumor growth ($p = 0.0082$).

The effects of MBP-vasostatin treatment on the human rhabdomyosarcoma cell line A-204 injected into nude mice were examined (Figure 13). Groups of athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (8×10^6 cell/mouse) s.c. with the human rhabdomyosarcoma cell line A-204. One day after cell injection, and continuing daily thereafter for 28 days, the mice received either control formulation buffer alone or MBP-vasostatin (100 µg/day, 6 days/week; Seq. I.D. No. 4). As shown, MBP-vasostatin reduced significantly the rate of growth of rhabdomyosarcoma in the mice ($p = 0.0006$). The mean tumor weight in the control group (2.449 g) was significantly greater ($p = 0.0009$) than the mean tumor weight in the treated group (0.859 g).

MBP-vasostatin (Seq. I.D. No. 4) was tested for its ability to reduce the growth of the human promyelomonocytic cell line HL60 inoculated s.c. into nude mice (Figure 14). Groups of athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (10×10^6 cell/mouse) with the human promyelomonocytic cell line HL60. One day after cell injection, and continuing daily thereafter for 24 days, the mice received either control formulation buffer alone or MBP-vasostatin (200 µg/day, 6 days/week; Seq. I.D. No. 4). The rate of tumor growth was significantly greater in the control group as compared to the treated group ($p < 0.001$). As shown, mice treated with MBP-vasostatin (Seq. I.D. No. 4) were significantly smaller than the mice treated with formulation buffer ($p < 0.001$).

The effect of MBP-vasostatin (Seq. I.D. No. 4) on human melanoma tumors transplanted into nude mice was also tested (Figure 15). Athymic BALB/c nude mice received 400 rad total body irradiation and 24 hours later were inoculated (7×10^6 cell/mouse) with the human melanoma cell line (A-375). Five days after cell injection, and continuing daily thereafter for 20 days, the mice received

either control formulation buffer alone or MBP-vasostatin (200 µg/day, 6 days/week; Seq. I.D. No. 4). As shown, MBP-vasostatin induced a significant reduction of Wilms tumor growth ($p=0.0059$).

These experiments indicate that the N-terminal 180 amino acid fragment of calreticulin, vasostatin (Seq. I.D. No. 4), can delay or prevent the growth of human tumors of various lineages established in nude mice.

VIII. A 61 Amino Acid Fragment of Calreticulin (amino acids 120-180; Seq. I.D. No. 5) Inhibits Tumor Growth *In Vivo*

In additional experiments, the 61 amino acid fragment of calreticulin encompassing amino acids 120-180 (Seq. I.D. No. 5) was tested for its ability to inhibit tumor growth. As described previously, this calreticulin fragment was active *in vitro* as an inhibitor of tumor growth and was active *in vivo* as an inhibitor of angiogenesis. To this end, groups of athymic mice were irradiated with 400 rad, and 24 hours later were injected s.c. with 4×10^6 cells from the Burkitt lymphoma cell line CA46. Beginning on the day of cell inoculation and continuing daily thereafter 6 days/week the mice received either control recombinant purified MBP (65 µg/day, 6 days/week), MBP-vasostatin (100 µg/day, 6 days/week; Seq. I.D. No. 4), or MBP-120-180 calreticulin (75 µg/day, 6 days/week; Seq. I.D. No. 5). Mice were injected for 42 days. As shown (Figure 16), the rate of tumor growth in the group of mice treated with MBP-vasostatin (Seq. I.D. No. 4) was significantly reduced when compared to controls ($p=0.0122$). In addition, the rate of tumor growth in the group of mice treated with MBP-120-180 calreticulin fragment (Seq. I.D. No. 5) was significantly reduced when compared to controls ($p=0.0058$). However, the rate of tumor growth in mice treated with MBP-vasostatin (Seq. I.D. No. 5) was not significantly different from the rate of tumor growth in the mice treated with the MBP-120-180 calreticulin fragment ($p=0.74$; Seq. I.D. No. 5). Hence, the 61 amino acid calreticulin fragment (Seq. I.D. No. 5) is active as an inhibitor of tumor growth *in vivo*.

IX. Antitumor Effects of Combination Therapy with Vasostatin and Interleukin-12

The effects of vasostatin and murine IL-12 (mIL-12), alone and together, were tested on human Burkitt lymphoma established in nude mice (Figure 17). In the first experiment (Figure 17A), the Burkitt cell line CA46 was inoculated s.c. into athymic mice, and 5 days later the established tumors were treated with daily s.c. inoculations (6 days/week) of either formulation buffer alone, MBP-vasostatin alone (100 µg/day; Seq. I.D. No. 4), mIL-12 alone (200 ng/day for 8 days followed by 100 ng/day), or the combination of MBP-vasostatin (Seq. I.D. No. 4) and mIL-12 (same dose used as single agents). All mice were sacrificed after 15 days of treatment. In a second experiment (Figure 17B), s.c. treatment with MBP-vasostatin (100 µg/day; 6 days/week; Seq. I.D. No. 4) or formulation buffer was started one day after the Burkitt Ca46 cells were injected into the nude mice, and s.c. treatment with IL-12 (100 ng/mouse; 6 days/week) was started 2 days later in mice that had received buffer alone or MBP-vasostatin (Seq. I.D. No. 4). The mice received 7 days treatment with MBP-vasostatin (Seq. I.D. No. 4) and 5 days treatment with IL-12, and were observed untreated for

14 days at which time all mice were sacrificed. The delayed addition of IL-12 and the shorter duration of IL-12 treatment in the second experiment resulted in IL-12 having a somewhat reduced effect on tumor growth and weight compared to the first experiment (Figures 17A and 17B).

However, in both experiments the rate of tumor growth and the mean weight of tumors was reduced by treatment with MBP-vasostatin (Seq. I.D. No. 4), mIL-12, or the combination of MBP-vasostatin (Seq. I.D. No. 4) plus mIL-12 compared with the control group treated with formulation buffer. Furthermore, the combination of IL-12 and MBP-vasostatin (Seq. I.D. No. 4) was more effective than either IL-12 alone ($p=0.018$ and $p<0.0001$, Figures 17A and 17B respectively) or MBP-vasostatin (Seq. I.D. No. 4) alone ($p=0.0029$ and $p=0.0046$, Figures 1A and 1B, respectively) at reducing the rate of tumor growth.

X. Therapeutic Activity Does not Stem From Binding to the Amino Acid Sequence KXaa₁FFXaa₂R (Seq. I.D. No. 11), or KXaa₁GFFKR (Seq. I.D. No. 10)

Attempts to identify a putative receptor for calreticulin and active calreticulin fragments on endothelial cells have focused initially on the possibility that such receptor might share the sequence motif KXaa₁FFXaa₂R (Seq. I.D. No. 11) present on distinct proteins known to bind calreticulin (Michalak et al. *Biochem. J.* 285:681-692, 1992; Coppolino et al., *Nature* 386:843-847, 1997; Coppolino et al., *J. Biol. Chem.* 270:23132-23138, 1995; Dedhar et al., *Nature* 367:480-483, 1994). Proteins known to bind calreticulin through a virtually identical amino acid sequence motif include the cytoplasmic domain of the alpha subunit of integrins (KXaa₁GFFKR; Seq. I.D. No. 10) and a family of steroid receptors (KXaa₁FFXaa₂R; Seq. I.D. No. 11). Specific examples of integrin amino acid sequences and steroid receptor sequences are provided in Table 5.

Table 5

Conservation of an Amino Acid Sequence Motif in the Integrin Alpha-subunit
Cytoplasmic Domains and in the Steroid Hormone Receptor Family
(Table derived from U.S. Patent No. 5,854,202 to Dedhar)

Integrins			Steroid Nuclear Receptors		
consensus sequence	KX _{aa1} GFFKR	Seq. ID No. 10	consensus sequence	KX _{aa1} FFX _{aa2} R	Seq. ID No. 11
a1	KIGFFKR	Seq. ID No. 12	RARa	*ACEGCKGFFRRSVQK	Seq. ID No. 21
a2	KLGFFKR	Seq. ID No. 13	T ₃ Rb	TCEGCKGFFRRTIQK	Seq. ID No. 22
a3	KGGFFKR	Seq. ID No. 14	VDR	TCEGCKGFFRRSMKR	Seq. ID No. 23
a4	KAGFFKR	Seq. ID No. 15	GR	TCGSCKVFFKRAVEG	Seq. ID No. 24
a5	KLGFFKR	Seq. ID No. 13	MR	TCGSCKVFFKRAVEG	Seq. ID No. 24
a6(A)	KCGFFKR	Seq. ID No. 17	AR	*TCGSCKVFFKRAAEK	Seq. ID No. 25
a6(B)	KCGFFKR	Seq. ID No. 16	PR	TCGSCKVFFKRAMEG	Seq. ID No. 26
a7	KLGFFKR	Seq. ID No. 13	ER	SCEGCKAFFKRSIQG	Seq. ID No. 27
a8	KCGFFDR	Seq. ID No. 17	RXR	SCEGCKGFFKRTVRK	Seq. ID No. 28
(chick)					
av	RMGFFKR	Seq. ID No. 18	Steroid Receptor TR2	TCEGCTGFFKRSIRK	Seq. ID No. 29
Mac-1	KLGFFKR	Seq. ID No. 13	Nerve growth factor induced protein 1-B	TCEGCKGFFKRTVQK	Seq. ID No. 30
p150	KVGFFKR	Seq. ID No. 19	Early response protein NAK1	TCEGCKGFFKRTVQK	Seq. ID No. 30
PS2	KCGFFNR	Seq. ID No. 20	Chorion Factor 1	SCEGCKGFFKRTVRK	Seq. ID No. 31
(Drosophila)					

*Sequences modified to conform to currently available sequence information.

To address this possibility, natural calreticulin fragments (purified from the culture supernatant of the VDS-O cell line) that were biologically active as inhibitors of endothelial cell growth, were tested to determine if they were capable of binding to amino acid sequences from the steroid nuclear receptor and the cytoplasmic domain of alpha integrin. If binding to these sequences is critical to the endothelial cell inhibitory function of calreticulin and its fragments, it should be possible to remove the biologically active molecules through affinity binding.

To this end, purified preparations of the VDS-O cell line containing full length calreticulin (Seq. I.D. No. 3) as well as N-terminal calreticulin fragments were prepared and tested to determine

how these components were retained over an affinity column. To prepare the affinity column, three peptides were synthesized (KLGFFRR, Seq. I.D. 13; KAFFKR, Seq. I.D. No. 33; and KVFFKR, Seq. I.D. No. 32), purified, and coupled to CNBr-activated Sepharose 4B (Pharmacia). First, the ability of calreticulin, purified from the culture supernatant of the lymphoblastoid cell line VDS-O, to
5 selectively bind to the column and be eluted by a buffer solution containing 20 mM EDTA and 100 mM NaCl, was verified. The purified preparations of VDS-O cell line containing both full length as well as N-terminal calreticulin fragments were then loaded on the column (Starting material). As shown in Figure 18 full length calreticulin present in the material loaded was retained by the column and could be eluted with 20 mM EDTA, whereas the calreticulin fragments were found in the flow-
10 through. These calreticulin fragments were subsequently found to be biologically active. Thus, the binding to the peptide sequence KXaa₁FFXaa₂R (Seq. I.D. No. 11) or the sequence KXaa₁GFFKR (Seq. I.D. No. 10) is not a property shared by calreticulin fragments that are active as inhibitors of endothelial cells growth and, moreover, the results suggest that KXaa₁FFXaa₂R (Seq. I.D. No. 11) and KXaa₁GFFKR (Seq. I.D. No. 10) are not part of a calreticulin receptor on endothelial cells.

15 To address this issue further, recombinant MBP-calreticulin fragment 120-180 (Seq. I.D. No. 5) was tested for its ability to bind to the affinity column described *supra*. Unlike MBP-calreticulin (Seq. I.D. No. 3) that bound to the column, the N-terminal calreticulin fragment (Seq. I.D. No. 5) failed to bind and was recovered in the flow-through.

The foregoing sections describe how the three newly discovered activities of calreticulin and calreticulin fragments were identified. The following Examples provide further guidance to one of
20 skill in the art with respect to the production of calreticulin and calreticulin fragments and variant proteins, as well as the pharmaceutical formulation and administration of these proteins.

Examples

25 Example 1. Expression and Purification of Calreticulin and Calreticulin Fragments

Calreticulin may be purified from the supernatant of Epstein-Barr virus (EBV)-immortalized cell lines as described above. Calreticulin may also be purified from a tissue source using
30 conventional biochemical techniques, or produced recombinantly in either prokaryotic or eukaryotic cells using methods well-known in the art (for example, those described in Sambrook et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor, New York, 1989). The recombinant expression of calreticulin and the 180 amino acid N-terminal fragment are described in Singh et al. (*Proc. Natl. Acad. Sci. U.S.A.* 91:12770-12774, 1994) and Atreya et al. (*J. Virol.* 69:3848-3851, 1995), respectively. Furthermore, the nucleic acid sequences encoding calreticulin are available on
35 GenBank, and include the cDNA sequence shown in Seq. I.D. No. 1.

Recombinant expression of calreticulin and calreticulin fragments may be conveniently obtained using commercial systems designed for optimal expression and purification of fusion proteins. Such fusion proteins typically include a protein tag that facilitates purification. Examples

of such systems include: the pMAL protein fusion and purification system (New England Biolabs, Inc., Beverly, MA); the GST gene fusion system (Amersham Pharmacia Biotech, Inc., Piscataway, NJ); and the pTrcHis expression vector system (Invitrogen, Carlsbad, CA). For example, the pMAL expression system utilizes a vector that adds a maltose binding protein to the expressed protein. The fusion protein is expressed in *E. coli*. and the fusion protein is purified from a crude cell extract using an amylose column. If necessary, the maltose binding protein domain can be cleaved from the fusion protein by treatment with a suitable protease, such as Factor Xa. The maltose binding fragment can then be removed from the preparation by passage over a second amylose column. Eukaryotic expression systems may also be employed, including *Pichia*, tobacco and *Baculovirus* expression systems, such as those available commercially from Invitrogen.

For each of these systems, the entire calreticulin protein may be produced by ligating the open reading frame (ORF) of calreticulin into the vector. To ensure effective expression, the ORF must be operably linked to the vector, i.e., must be joined such that the reading frame of the ORF is aligned with the reading frame of the protein tag. Where fragments of calreticulin are to be expressed, an ORF encoding the desired fragment may be amplified by polymerase chain reaction (PCR) from the calreticulin cDNA, cloned, purified and then ligated into the expression vector. Alternatively, the amplified fragment may be ligated directly into the expression vector. It may also be possible, depending on the availability of suitable restriction sites in the calreticulin cDNA to obtain the desired fragment by appropriate restriction endonuclease digestion, such that it can be directly cloned into the expression vector.

Purification of the expressed protein can be achieved either using the purification regimen appropriate for the expression tag (if a commercial expression/purification system is used), or conventional affinity chromatography using antibodies, preferably monoclonal antibodies, that recognize the appropriate regions of calreticulin may be employed.

Where calreticulin fragments are to be used, such fragments may alternatively be generated through digestion of the full-length calreticulin protein with various proteases. The fragments may then be separated based on their unique size, charge or other characteristics. Calreticulin fragments may also be synthetically generated through the use of known peptide synthesis methods.

Example 2. Assessing Therapeutic Activity

Following the purification of calreticulin or a fragment of calreticulin, the biological activity can be assessed using the methods described above. Specifically, the bFGF-induced endothelial cell proliferation assay can be used to determine endothelial cell inhibition, the Matrigel assay can be used to measure the inhibition of angiogenesis, and the athymic mouse/human Burkitt lymphomas model can be used to quantitate tumor inhibition.

When testing a calreticulin or calreticulin fragment preparation for activity, the bFGF-induced endothelial cell proliferation assay is generally first used. One skilled in the art will appreciate that this assay yields relatively quick results at relatively low cost.

Example 3. Calreticulin Sequence Variants

While the amino acid sequence of the prototypical human calreticulin protein is provided in Seq. I.D. No. 2, and the sequence of a cDNA molecule encoding this protein is given in Seq. I.D. No. 6, one of skill in the art will appreciate that the practice of this invention is not limited to these
5 precise sequences. Thus, the invention may be practiced with molecules that differ from the exact molecules disclosed, but which retain the requisite biological activity.

Furthermore, variants of calreticulin and fragments of calreticulin that have been modified such that they do not bind to the amino acid motif described *supra* are of particular interest. These variants will retain the ability to specifically bind to endothelial cells, but will not be capable of
10 binding the amino acid sequences shown in Seq. I.D. Nos. 11 and/or 10. This amino acid sequence motif is found in a family of steroid receptors (glucocorticoid, mineralcorticoid, progesterone and androgen receptors: KVFFKR, Seq. I.D. No. 32; estrogen receptor: KAFFKR, Seq. I.D. No. 33; thyroid hormone receptor: KSFFRR, Seq. I.D. No. 34; and retinoic acid receptor: KGFFRR, Seq. I.D. 35). Furthermore, it is also likely that therapeutically active variants and fragments of calreticulin do
15 not bind to the consensus sequences shown in Seq. I.D. Nos. 10 and 11. A specific example of a naturally occurring sequence that falls into the integrin consensus sequence Seq. I.D. No. 10, is shown in Seq. I.D. No. 13. However, it is foreseeable that a calreticulin variant or fragment that does not bind to the consensus sequences shown in Seq. I.D. Nos. 10 and 11 may be used in combination with full length calreticulin or a derivative of calreticulin that does bind to Seq. I.D. Nos. 10 and 11.

As mentioned above, the fragments and variants of calreticulin described *supra*, are characterized by their ability to inhibit endothelial cell growth, angiogenesis, and/or tumor growth. These abilities, however, are further defined as at least 30% inhibition of endothelial cell growth, angiogenesis, and/or tumor growth. However, it is likely that some therapeutically active fragments and variants of calreticulin will show an increased level of one or more of these biological activities.
25 For example, some variants and fragments of calreticulin will show at least 40% inhibition, at least 50% inhibition, at least 60% inhibition, or at least 70% inhibition. Moreover, using the assays described above it is now possible to individually assess the biological activity of a given variant or fragment of calreticulin.

The therapeutically effective fragments and variants of calreticulin are also characterized by
30 the number of amino acid residues that they contain. For example, in some instances it may be desirable to use relatively short fragments and variants of calreticulin. These short fragments and variants of calreticulin may contain at least 5, 10, 20, or 30 contiguous amino acids residues of the calreticulin sequence. However, such short fragments and variants of calreticulin will maintain at least one of the biological activities described *supra*.

Additionally, it is possible to vary the cDNA sequences encoding therapeutically effective fragments or variants of calreticulin while still encoding a protein having the desired biological activity. In their simplest form, such sequence variants may differ from the disclosed sequences by alteration of the coding region to fit the codon usage bias of the particular organism into which the molecule is to be introduced. Additionally, the coding region may be altered by taking advantage of
35

the degeneracy of the genetic code to alter the coding sequence in such a way that, while the nucleotide sequence is substantially altered, it nevertheless encodes a protein having an amino acid sequence identical or substantially similar to the disclosed calreticulin protein sequence. For example, the first amino acid residue of the mature calreticulin protein is glutamic acid (Glu, E). This is encoded in the calreticulin open reading frame (ORF) by the nucleotide codon triplet GAG. Because of the degeneracy of the genetic code, one other nucleotide codon, GAA, also encodes for glutamic acid. Thus, the nucleotide sequence of the calreticulin ORF could be changed at this position to GAA without affecting the amino acid composition of the encoded protein or the characteristics of the protein.

As previously mentioned, the invention may also be practiced with calreticulin and calreticulin fragments that vary in amino acid sequence from the sequence shown in Seq. I.D. No. 2 (i.e. peptides that have been modified such that they do not bind to the consensus sequences described *supra* (Seq. I.D. Nos. 10 and 11). Variant calreticulin proteins include proteins that differ in amino acid sequence from the endogenous calreticulin sequence disclosed but which retain the specified biological activity. Such proteins may be produced by manipulating the nucleotide sequence of ORF that encodes the protein, for example by site-directed mutagenesis or the polymerase chain reaction. The simplest modifications involve the substitution of one or more amino acids for amino acids having similar biochemical properties. These so-called conservative substitutions are likely to have minimal impact on the activity of the resultant protein. Table 6 shows amino acids which may be substituted for an original amino acid in a protein and which are regarded as conservative substitutions.

Table 6.

Original Residue	Conservative Substitutions
Ala	ser
Asn	gln; his
Asp	glu
Cys	ser
Gln	asn
Glu	asp
Gly	pro
His	asn; gln
Ile	leu; val
Leu	ile; val
Lys	arg; gln; glu
Met	leu; ile
Phe	met; leu; tyr
Ser	thr
Thr	ser
Trp	tyr
Tyr	trp; phe
Val	ile; leu

More substantial changes in function or other features may be obtained by selecting substitutions that are less conservative than those in Table 4, i.e., selecting residues that differ more significantly in their effect on maintaining (a) the structure of the polypeptide backbone in the area of the substitution, for example, as a sheet or helical conformation, (b) the charge or hydrophobicity of the molecule at the target site, or (c) the bulk of the side chain. The substitutions which in general are expected to produce the greatest changes in protein properties will be those in which (a) a hydrophilic residue, e.g., seryl or threonyl, is substituted for (or by) a hydrophobic residue, e.g., leucyl, isoleucyl, phenylalanyl, valyl or alanyl; (b) a cysteine or proline is substituted for (or by) any other residue; (c) a residue having an electropositive side chain, e.g., lysyl, arginyl, or histadyl, is substituted for (or by) an electronegative residue, e.g., glutamyl or aspartyl; or (d) a residue having a bulky side chain, e.g., phenylalanine, is substituted for (or by) one not having a side chain, e.g., glycine. The effects of these amino acid substitutions or deletions or additions may be assessed through the use of the biological assays described above.

Example 4. Incorporation of Therapeutically Effective Fragments and Variants of Calreticulin into Pharmaceutical Compositions

For administration to animals, purified calreticulin or calreticulin fragments are generally combined with a pharmaceutically acceptable carrier. Pharmaceutical preparations may contain only calreticulin or a single calreticulin fragment, or may be composed of calreticulin combined with one or more calreticulin fragments, or may be composed of multiple calreticulin fragments. In general, the nature of the carrier will depend on the particular mode of administration being employed. For instance, parenteral formulations usually comprise injectable fluids that include pharmaceutically and physiologically acceptable fluids such as water, physiological saline, balanced salt solutions, aqueous dextrose, glycerol, human albumin or the like as a vehicle. For solid compositions (e.g., powder, pill, tablet, or capsule forms), conventional non-toxic solid carriers can include, for example,

pharmaceutical grades of mannitol, lactose, starch, or magnesium stearate. In addition to biologically-neutral carriers, pharmaceutical compositions to be administered can contain minor amounts of non-toxic auxiliary substances, such as wetting or emulsifying agents, preservatives, and pH buffering agents and the like, for example sodium acetate or sorbitan monolaurate.

5 As is known in the art, protein-based pharmaceuticals may be only inefficiently delivered through ingestion. However, pill-based forms of pharmaceutical proteins may be alternatively be administered subcutaneously, particularly if formulated in a slow-release composition. Slow-release formulations may be produced by combining the target protein with a biocompatible matrix, such as cholesterol. Another possible method of administering protein pharmaceuticals is through the use of
10 mini osmotic pumps. As stated above a biocompatible carrier would also be used in conjunction with this method of delivery.

It is also contemplated that calreticulin could be delivered to cells in the nucleic acid form and subsequently translated by the host cell. This could be done, for example through the use of viral vectors or liposomes. Liposomes could also be used for the delivery of the protein itself.

15 The therapeutically effective fragments may also be delivered in conjunction with other therapeutic agents. These additional therapeutic agents can be angiogenesis inhibitors such as, platelet-factor-4, IP-10 (interferon (IFN)- γ inducible protein-10), MIG (Monokine induced by IFN- γ), IFN- α , angiostatin, endostatin, fumagillin, AGM-1470, thrombospondin, a fragment of prolactin, antibody against the integrin $\alpha_v\beta_3$, IL-12, cleaved conformation of the serpin antithrombin (O'Reilly
20 *et al.*, *Science* 285:1926, 1999), thalidomide, and mixtures thereof. The additional therapeutics could also be chemotherapeutics, hormones, anti-inflammatory agents, antibiotics, radiation therapy and/or anti-viral agents.

The pharmaceutical compositions of the present invention may be administered by any means that achieve their intended purpose. Amounts and regimens for the administration of
25 calreticulin, or a therapeutically active fragment thereof, can be determined readily by those with ordinary skill in the clinical art of treating diseases associated with angiogenesis, endothelial cell growth and/or tumor growth. For use in treating these conditions, the described proteins are administered in an amount effective to inhibit angiogenesis, endothelial cell growth and/or tumor growth. The peptides or proteins may be administered to a host *in vivo*, such as for example, through
30 systemic administration, such as intravenous or intraperitoneal administration. Also, the peptides or proteins may be administered intralesionally: i.e. the peptide or protein is injected directly into the tumor or affected area.

Effective doses of calreticulin and calreticulin fragments for therapeutic application will vary depending on the nature and severity of the condition to be treated, the age and condition of the
35 subject and other clinical factors. Thus, the final determination of the appropriate treatment regimen will be made by the attending clinician. Typically, the dose range will be from about 0.1 $\mu\text{g/kg}$ body weight to about 100 mg/kg body weight. Other suitable ranges include doses of from about 1 $\mu\text{g/kg}$ to 10 mg/kg body weight. The dosing schedule may vary from once a week to daily depending on a

number of clinical factors, such as the subject's sensitivity to the protein. Examples of dosing schedules are 3 µg/kg administered twice a week, three times a week or daily; a dose of 7 µg/kg twice a week, three times a week or daily; a dose of 10 µg/kg twice a week, three times a week or daily; or a dose of 30 µg/kg twice a week, three times a week or daily. In the case of a more aggressive disease it may be preferable to administer doses such as those described above by alternate routes including intravenously or intrathecally. Continuous infusion may also be appropriate.

As mentioned above calreticulin and calreticulin fragments will be useful for the treatment of diseases associated with unwanted angiogenesis. Angiogenesis is commonly associated with ocular diseases. Some of these diseases are, retrolental fibroplasia, trachoma, retinal neovascularization, macular degeneration, diabetic retinopathy and neovascular glaucoma, corneal graft rejection and contact lens overwear. Other non-ocular diseases, which are associated with unwanted angiogenesis, can also be treated with calreticulin and fragments of calreticulin. Examples of such diseases are, periodontal disease, psoriasis, angiofibromas, immune-inflammation, atherosclerosis, excessive wound repair, non-immune inflammation, Crohn's disease, and systemic lupus. Examples of diseases that are associated with immune-inflammation are rheumatoid arthritis, systemic lupus erythematosus, thyroiditis, Goodpasture's Syndrome, systemic vasculitis, scleroderma, Sjogren's syndrome, sarcoidosis or primary biliary cirrhosis.

Calreticulin and fragments of calreticulin may also be useful for the treatment diseases with unknown etiology, for example, Kaposi's sarcoma.

In addition to being used for the treatment of disease, calreticulin and fragments of calreticulin, may be useful as preventative agents. For instance, calreticulin may be useful as a preventative of pregnancy by inhibiting angiogenesis at the site of implantation. Calreticulin and the effective fragments might also be used as a preventative of injury from radiation and/or chemotherapy. Used in this way the therapeutically active fragments or variants would be administered prior to the radiation and/or chemotherapy treatment and the resulting inhibition of angiogenesis would serve to protect hematopoietic cells from injury.

Calreticulin and calreticulin fragments can also be used to treat a variety of malignancies and related disorders, such as leukemias, including acute leukemias (such as acute lymphocytic leukemia, acute myelocytic leukemia, and myeloblastic, promyelocytic, myelomonocytic, monocytic and erythroleukemia), chronic leukemias (such as chronic myelocytic (granulocytic) leukemia and chronic lymphocytic leukemia), polycythemia vera, lymphoma, Hodgkin's disease, non-Hodgkin's lymphoma, multiple myeloma, Waldenström's macroglobulinemia, heavy chain disease), as well as solid tumors such as sarcomas and carcinomas. fibrosarcoma, myxosarcoma, liposarcoma, chondrosarcoma, osteogenic sarcoma, and other sarcomas, synovioma, mesothelioma, Ewing's tumor, leiomyosarcoma, rhabdomyosarcoma, colon carcinoma, pancreatic cancer, breast cancer, ovarian cancer, prostate cancer, squamous cell carcinoma, basal cell carcinoma, adenocarcinoma, sweat gland carcinoma, sebaceous gland carcinoma, papillary carcinoma, papillary adenocarcinomas, medullary carcinoma, bronchogenic carcinoma, renal cell carcinoma, hepatoma, bile duct carcinoma,

choriocarcinoma, Wilms' tumor, cervical cancer, testicular tumor, bladder carcinoma, CNS tumors (such as a glioma, astrocytoma, medulloblastoma, craniopharyngioma, ependymoma, pinealoma, hemangioblastoma, acoustic neuroma, oligodendroglioma, meningioma, melanoma, neuroblastoma and retinoblastoma). As stated above the precise details of the clinical treatment will depend on the
5 specific tumor type and the severity of the condition.

Furthermore, calreticulin and fragments of calreticulin have been shown to inhibit angiogenesis that is induced by basic fibroblast growth factor (see, endothelial cell proliferation assay, discussed above). Therefore, one of ordinary skill will appreciate that calreticulin and fragments of calreticulin will also inhibit angiogenesis, which is induced by other compounds. These
10 angiogenesis inducing compounds could be for example, acidic fibroblast growth factor, Vascular Endothelial Growth Factor (VEGF), hepatocyte growth factor, Interleukin (IL)-15, IL-12, IL-8, platelet-derived endothelial cell growth factor (PDEC GF), angiogenin, Cripto (Ebert *et al.*, *Cancer Res.*, 59:4502-4505, 1999), Transforming Growth Factor (TGF)- β , Tumor Necrosis Factor (TNF) α , and angiogenin.

15 Having illustrated and described the principles of the invention in multiple embodiments and examples, it should be apparent to those skilled in the art that the invention can be modified in arrangement and detail without departing from such principles. We claim all modifications coming within the spirit and scope of the following claims.

Claims

We claim:

- 5 1. A method of inhibiting endothelial cell growth, comprising contacting endothelial cells with a pharmaceutical composition comprising at least one protein selected from the group consisting of:
- (a) therapeutically effective fragments of calreticulin;
 - (b) therapeutically effective variants of calreticulin;
 - (c) calreticulin; and
 - (d) therapeutically effective variants of the fragments of (a).
- 10 2. The method of claim 1 wherein the pharmaceutical composition further comprises a pharmaceutically acceptable carrier.
- 15 3. The method of claim 1 wherein the therapeutically effective fragment of calreticulin comprises a polypeptide of about 180 amino acids from the N-terminus of calreticulin.
4. The method of claim 1 wherein the therapeutically effective fragment of calreticulin consists essentially of an amino acid sequence selected from the group consisting of:
- (a) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 3;
 - 20 (b) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 4;
 - (c) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 5;
 - (d) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 6;
 - 25 (e) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 8; and
 - (f) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 9.
- 30 5. A method of inhibiting angiogenesis in a subject, comprising administering to the subject an effective amount of a pharmaceutical composition comprising at least one protein selected from the group consisting of:
- 35 (a) therapeutically effective fragments of calreticulin;
 - (b) therapeutically effective variants of calreticulin;
 - (c) calreticulin; and
 - (d) therapeutically effective variants of the fragments of (a).

6. The method of claim 5 wherein the pharmaceutical composition further comprises a pharmaceutically acceptable carrier.

7. The method of claim 5 wherein the therapeutically effective fragment of calreticulin comprises a polypeptide of about 180 amino acids from the N-terminus of calreticulin.

8. The method of claim 5 wherein the therapeutically effective fragment of calreticulin consists essentially of an amino acid sequence selected from the group consisting of:

(a) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 3;

(b) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 4;

(c) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 5;

(d) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 6;

(e) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 8; and

f) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No.

9.

9. The method of claim 5 wherein the angiogenesis is associated with a disease, other than a tumor, that is associated with neovascularization.

10. The method of claim 9 wherein angiogenesis is inhibited in a disease selected from a group consisting of diabetic retinopathy, retrolental fibroplasia, trachoma, neovascular glaucoma, psoriasis, angiofibromas, immune-inflammation, atherosclerosis, excessive wound repair, retinal neovascularization, macular degeneration, corneal graft rejection, contact lens overwear, Crohn's disease and non-immune inflammation.

11. The method of claim 9 wherein the disease is selected from a group consisting of rheumatoid arthritis, systemic lupus erythematosus, thyroiditis, Goodpasture's Syndrome, systemic vasculitis, scleroderma, Sjogren's syndrome, sarcoidosis and primary biliary cirrhosis.

12. A method of treatment of Kaposi's sarcoma, comprising administering to the subject an effective amount of pharmaceutical composition comprising at least one protein selected from the group consisting of:

(a) therapeutically effective fragments of calreticulin;

(b) therapeutically effective variants of calreticulin;

- (c) calreticulin; and
- (d) therapeutically effective variants of the fragments of (a).

- 5 13. The method of claim 5 further comprising providing a second anti-angiogenic agent selected from a group consisting of platelet-factor-4, IP-10 (interferon (IFN)- γ inducible protein-10), MIG (Monokine induced by IFN- γ), INF- γ , IFN- α , angiostatin, endostatin, fumagillin, AGM-1470, thrombospondin, a fragment of prolactin, antibody against the integrin $\alpha_v\beta_3$, IL-12, cleaved conformation of the serpin antithrombin, thalidomide, and mixtures thereof.
- 10 14. The method of claim 5 further comprising administering a chemotherapeutic agent.
- 15 15. The method of claim 5 further comprising administering a hormone.
- 16 16. The method of claim 5 further comprising administering an anti-inflammatory agent.
- 17 17. The method of claim 5 further comprising administering an anti-viral agent.
- 18 18. The method of claim 5 wherein the angiogenesis is inhibited in pregnancy.
- 20 19. The method of claim 5 wherein the angiogenesis is inhibited to terminate pregnancy.
- 20 20. The method of claim 5 wherein the angiogenesis is inhibited in periodontal disease.
- 25 21. The method of claim 20 further comprising administering an antibiotic.
- 25 22. The method of claim 5 wherein the angiogenesis is inhibited in radiation induced injury.
- 30 23. The method of claim 5 wherein the angiogenesis is inhibited in chemotherapy induced injury.
- 35 24. The method of claim 5 wherein the pharmaceutical composition inhibits angiogenesis which is stimulated by an angiogenesis inducer selected from a group consisting of, basic fibroblast growth factor, acidic fibroblast growth factor, Vascular Endothelial Growth Factor (VEGF), hepatocyte growth factor, Interleukin (IL)-15, IL-8, platelet-derived endothelial cell growth factor (PDECGF), Transforming Growth Factor (TGF)- β , Tumor necrosis Factor (TNF) α , angiogenin, Cripto, and mixtures thereof.
25. The method of claim 5, wherein the subject is immunocompromized due to T-lymphocyte deficiency.

26. A method of inhibiting tumor growth, comprising contacting tumor cells with an effective amount of a pharmaceutical composition comprising at least one protein selected from the group consisting of:

- (a) therapeutically effective fragments of calreticulin;
- (b) therapeutically effective variants of calreticulin;
- (c) calreticulin; and
- (d) therapeutically effective variants of the fragments of (a).

27. The method of claim 26 wherein the pharmaceutical composition further comprises a pharmaceutically acceptable carrier.

28. The method of claim 26 wherein the therapeutically effective fragment of calreticulin comprises a polypeptide of about 180 amino acids from the N-terminus of calreticulin.

29. The method of claim 26 wherein the therapeutically effective fragment of calreticulin consists essentially of an amino acid sequence selected from the group consisting of:

- (a) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 4;
- (b) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 5;
- (c) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 6;
- (d) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 8; and
- (e) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 9.

30. The method of claim 26 wherein the tumor growth occurs in a subject and is inhibited by administering to the subject an effective amount of the pharmaceutical composition.

31. A protein consisting essentially of an amino acid sequence selected from the group consisting of:

- (a) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 4;
- (b) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 5;
- (c) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 6;

(d) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 8; and

(e) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 9.

32. A composition comprising a protein according to claim 31, and a pharmaceutically acceptable carrier.

33. A vector comprising a nucleotide sequence encoding a protein with an amino acid sequence selected from the group consisting of:

(a) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 3;

(b) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 4;

(c) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 5;

(d) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 6;

(e) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D.

No. 8; and

f) at least 5 contiguous amino acid residues of the sequence shown in Seq. I.D. No. 9.

34. A host cell comprising the vector according to claim 33.

35. A method of isolating a therapeutically effective fragment of calreticulin from a sample comprising:

(a) contacting the sample with at least one peptide where the peptide has the amino acid sequence selected from the group consisting of Seq. I.D. No. 10 and Seq. I.D. No. 11;

(b) recovering the portion of the sample that does not bind to the peptide; and

(c) assaying the recovered portion of the sample for therapeutic activity.

36. A therapeutically effective fragment of calreticulin identified by the method of claim 35.

37. A vector comprising a nucleotide sequence encoding the therapeutically effective fragment of claim 36.

38. A host cell comprising the vector according to claim 37.

39. A method of isolating a therapeutically effective variant of calreticulin from a sample comprising:

- (a) contacting the sample with at least one peptide where the peptide has the amino acid sequence selected from the group consisting of Seq. I.D. No. 10 and Seq. I.D. No. 11;
- (b) recovering the portion of the sample that does not bind to the peptide; and
- (c) assaying the recovered portion of the sample for therapeutic activity.

40. A therapeutically effective variant of calreticulin identified by the method of claim 39.

41. A vector comprising a nucleotide sequence encoding the therapeutically effective variant of claim 40.

42. A host cell comprising the vector according to claim 41.

43. A therapeutically effective fragment of calreticulin that:

- (a) does not bind to the amino acid sequence shown in Seq. I.D. No. 10; and
- (b) displays a biological activity selected from the group consisting of: at least 30% inhibition of angiogenesis, at least 30% inhibition of tumor growth, and at least 30% inhibition of endothelial cell growth.

44. The therapeutically effective fragment of claim 43, wherein the biological activity is selected from the group consisting of: at least 40% inhibition of angiogenesis, at least 40% inhibition of tumor growth, and at least 40% inhibition of endothelial cell growth.

45. A vector comprising a nucleotide sequence encoding the therapeutically effective fragment of claim 43.

46. A host cell comprising the vector according to claim 45.

47. A therapeutically effective variant of calreticulin that:

- (a) does not bind to the amino acid sequence shown in Seq. I.D. No. 10; and
- (b) displays a biological activity selected from the group consisting of: at least 30% inhibition of angiogenesis, at least 30% inhibition of tumor growth, and at least 30% inhibition of endothelial cell growth.

48. The therapeutically effective fragment of claim 42, wherein the biological activity is selected from the group consisting of: at least 40% inhibition of angiogenesis, at least 40% inhibition of tumor growth, and at least 40% inhibition of endothelial cell growth.

49. A vector comprising a nucleotide sequence encoding the therapeutically effective variant of claim 47.

50. A host cell comprising the vector according to claim 49.

51. A mimetic of the protein of claim 31.

52. A mimetic of the therapeutically effective fragment of claim 36.

53. A mimetic of the therapeutically effective variant of claim 40.

54. A mimetic of the therapeutically effective fragment of claim 43.

55. A mimetic of the therapeutically effective variant of claim 47.

56. The method of claim 5 further comprising administering radiation therapy.

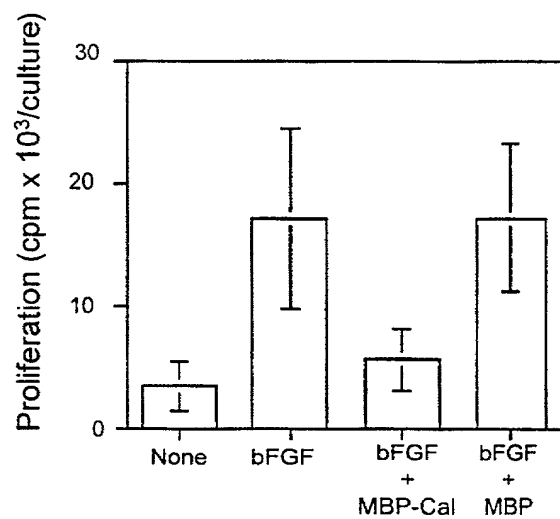


Fig. 1

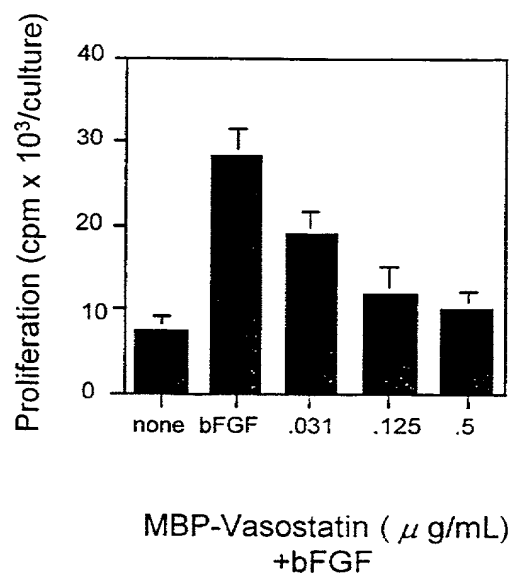


Fig. 2

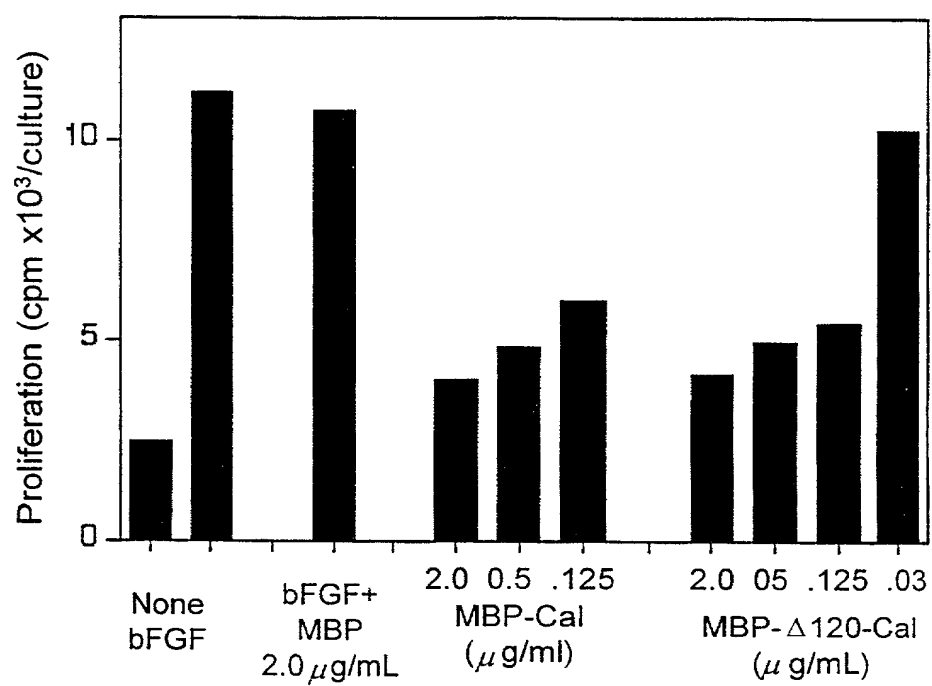


Fig. 3

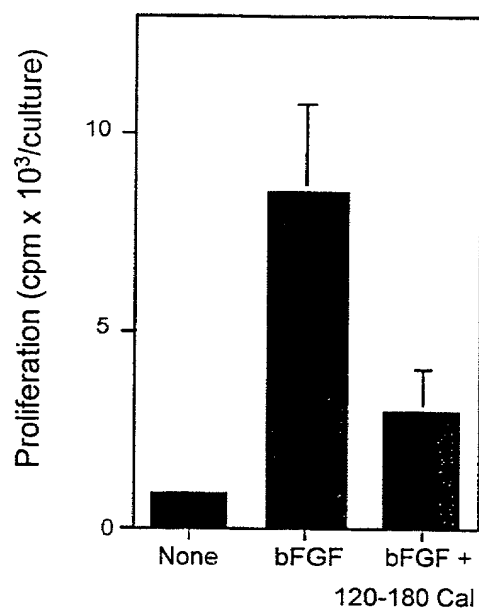


Fig. 4

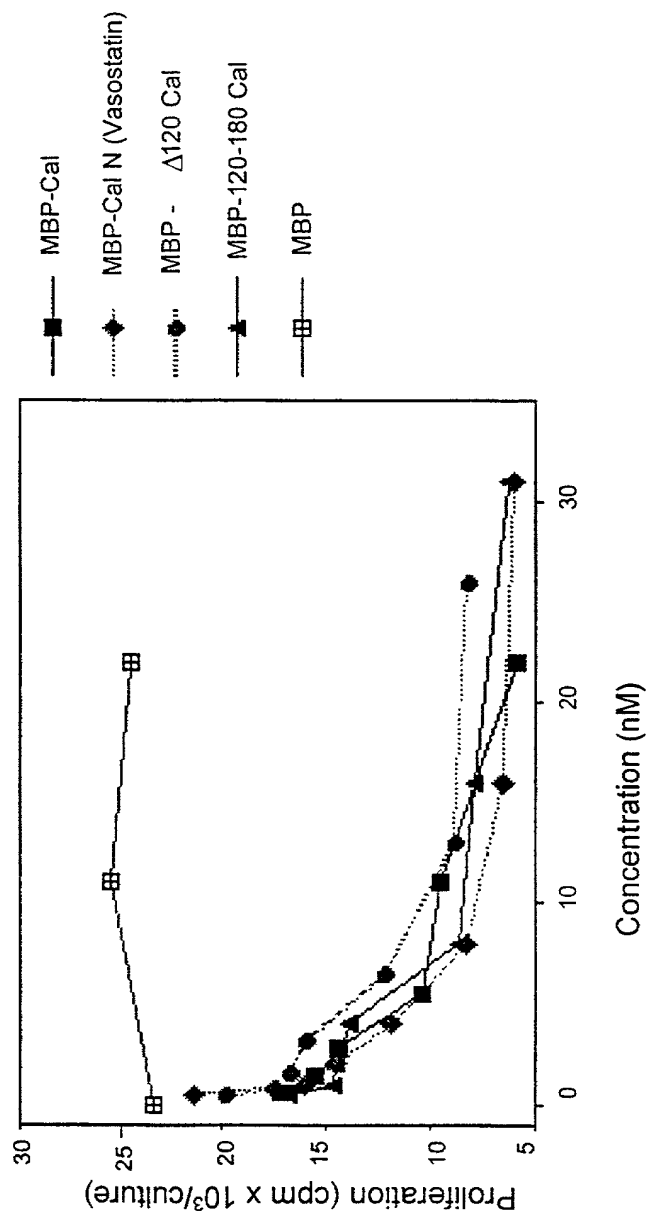


Fig. 5

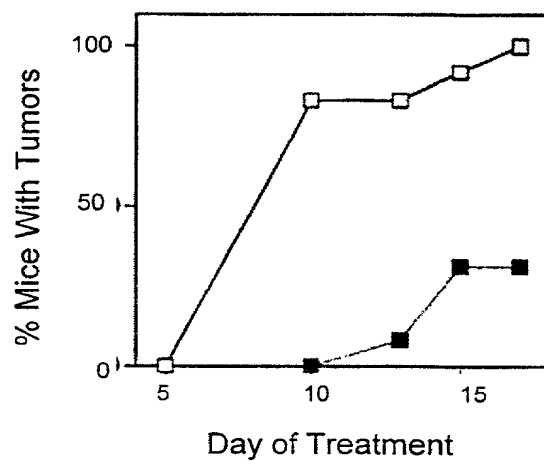


Fig. 6A

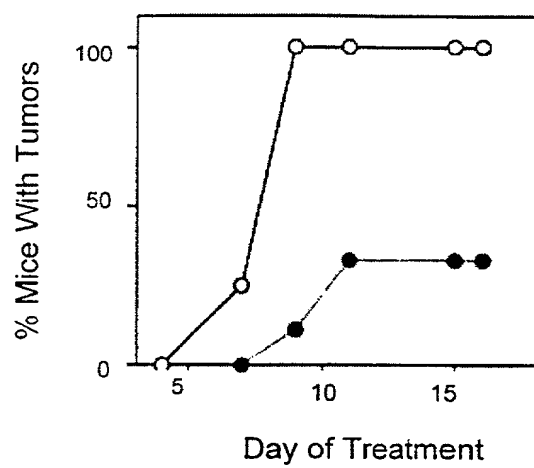


Fig. 6B

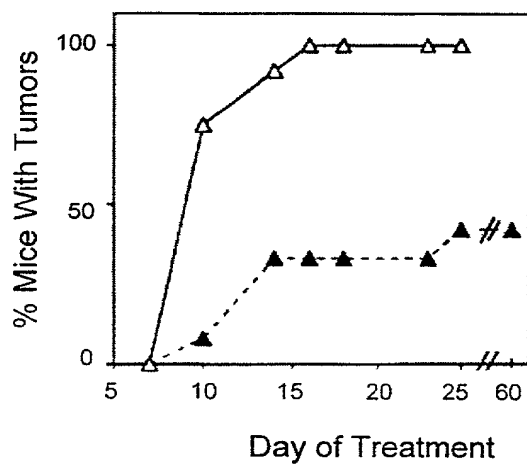


Fig. 6C

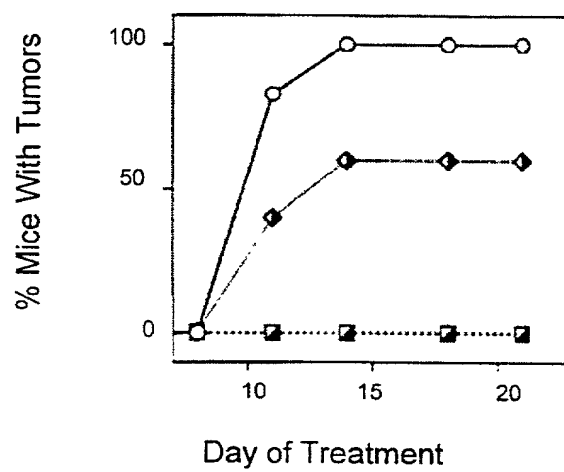


Fig. 6D

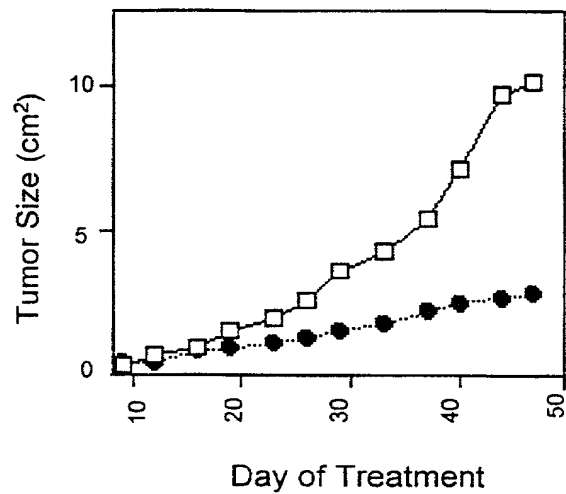


Fig. 7A

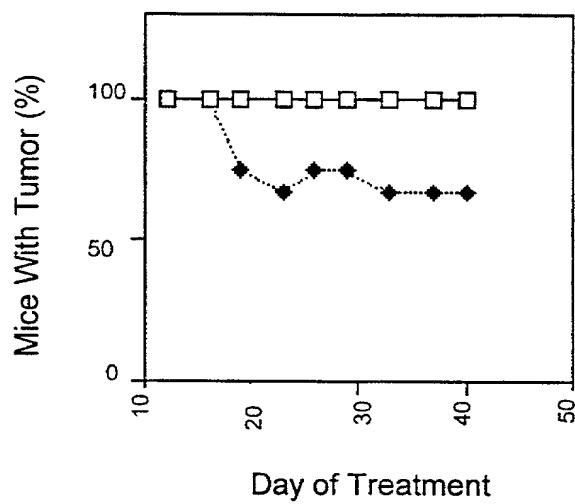


Fig. 7B

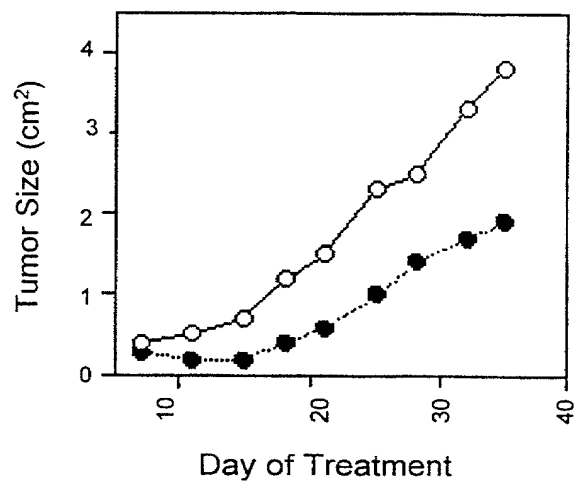


Fig. 8

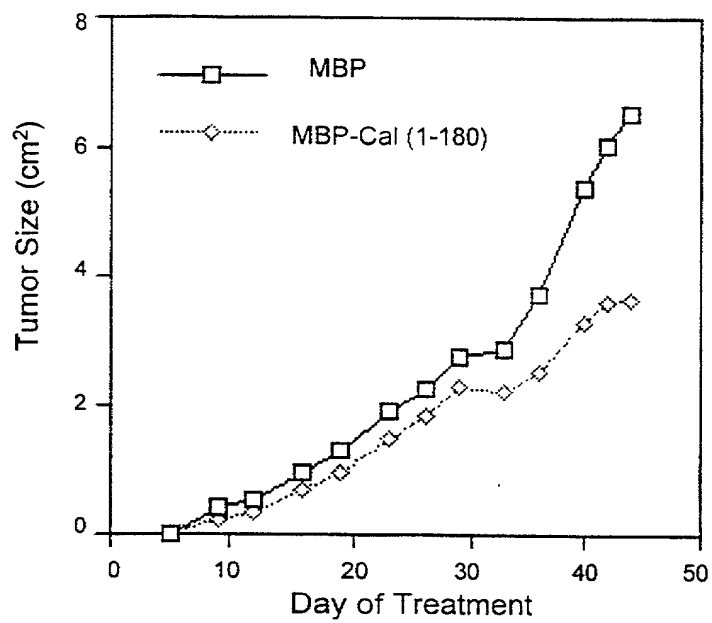


Fig. 9A

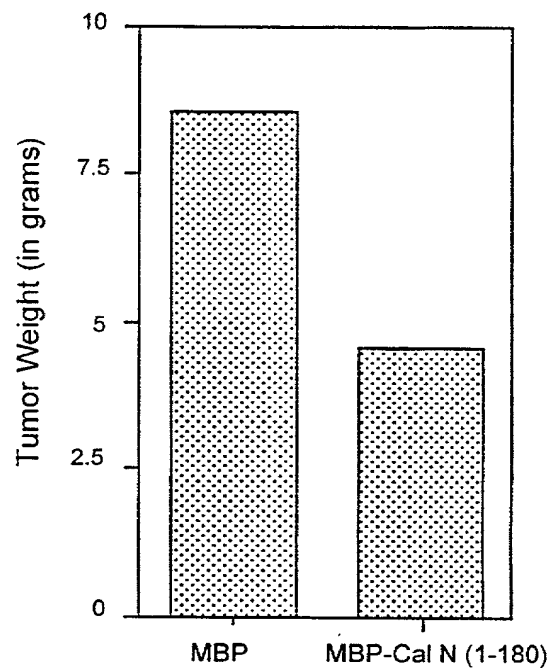


Fig. 9B

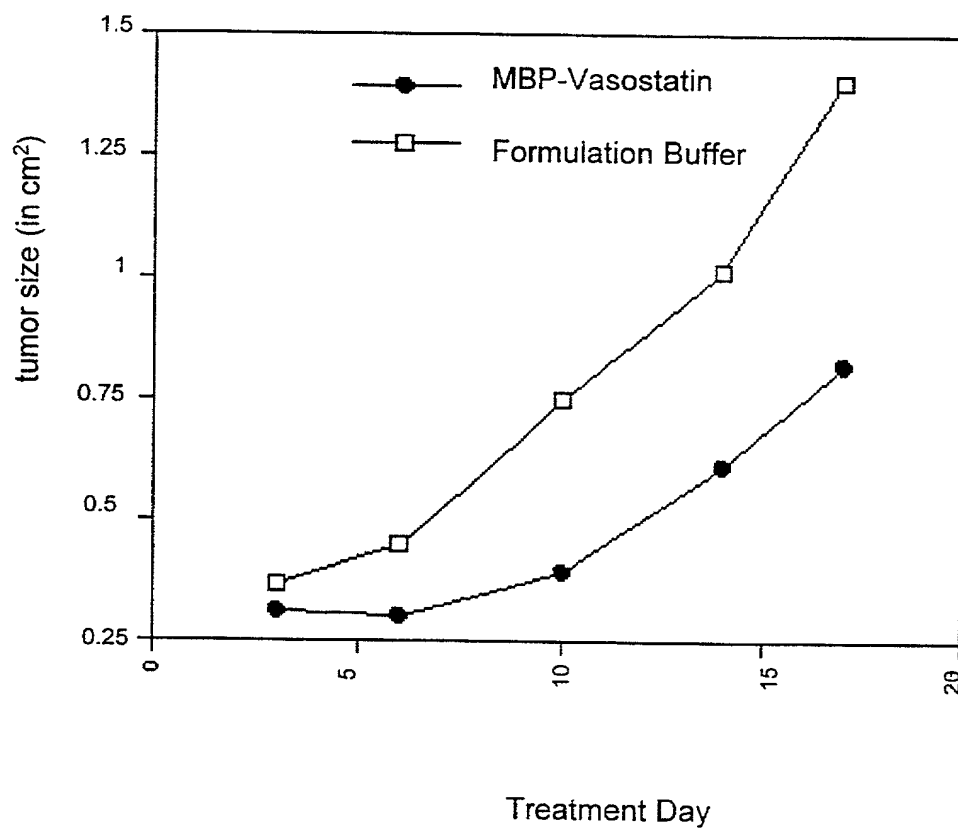


Fig. 10

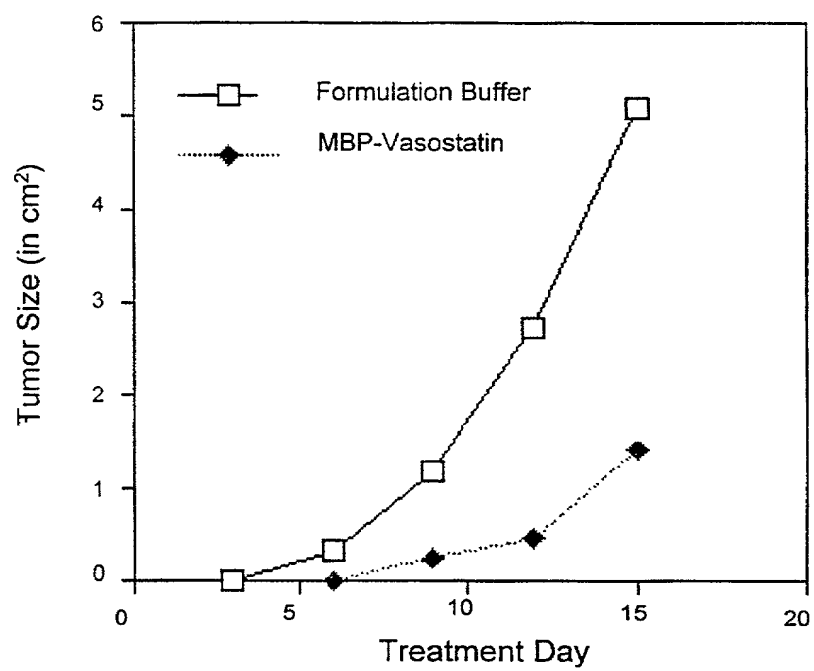


Fig. 11A

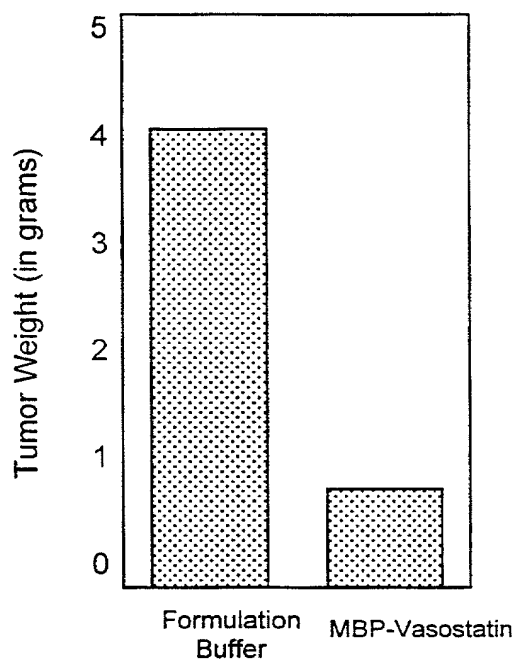


Fig. 11B

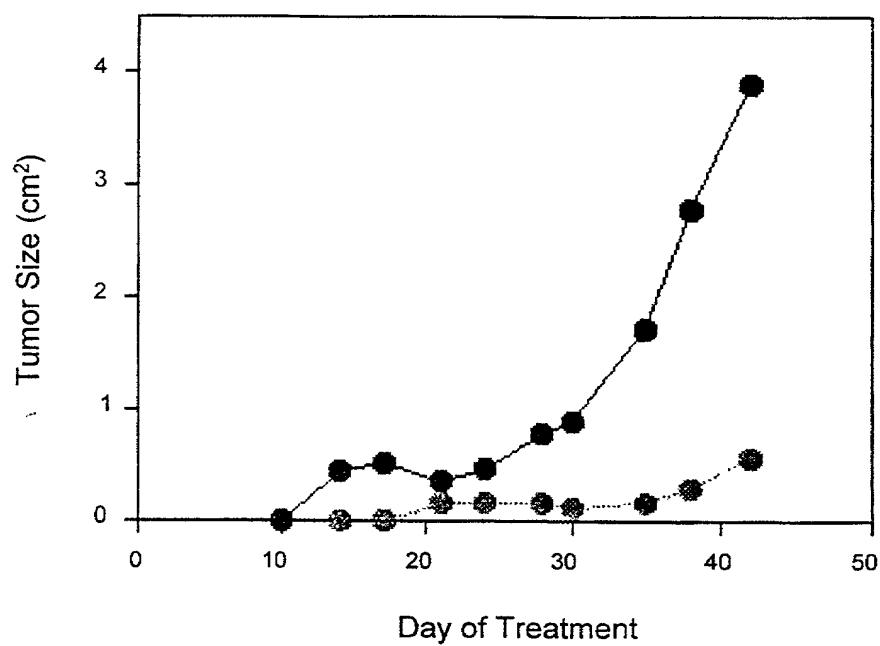


Fig. 12

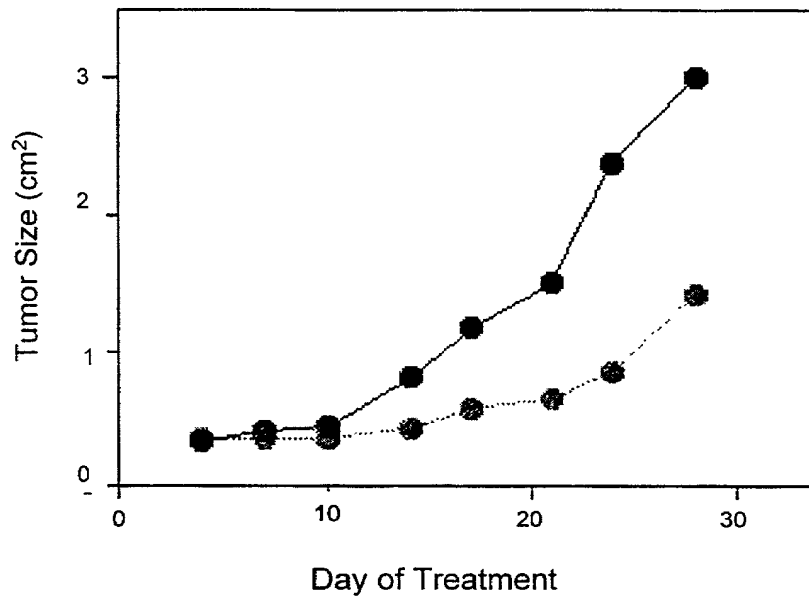


Fig. 13

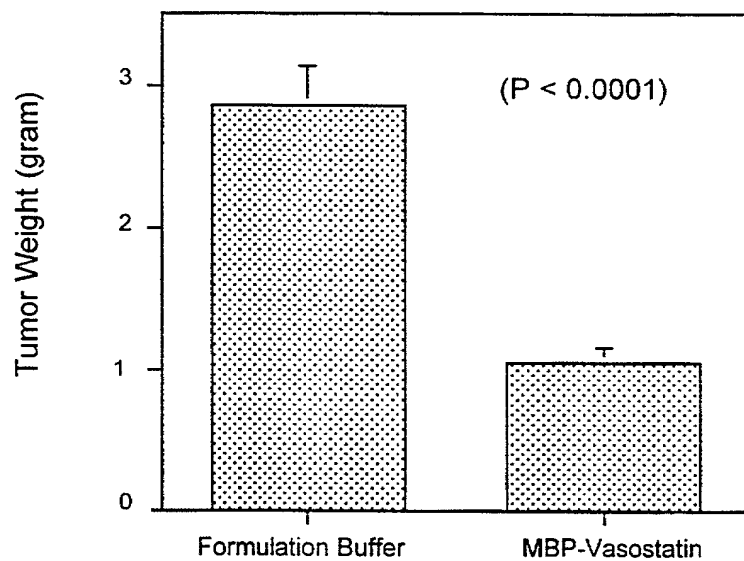


Fig. 14

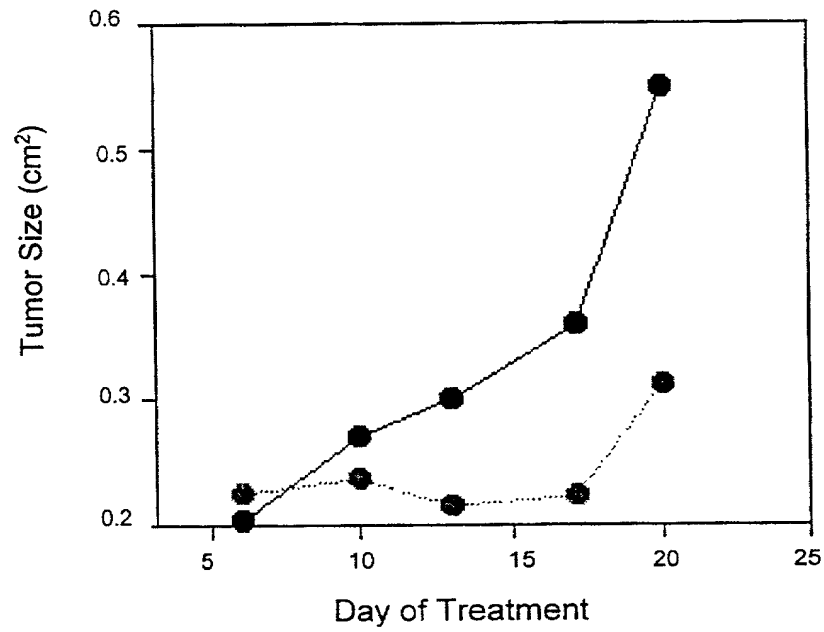


Fig. 15

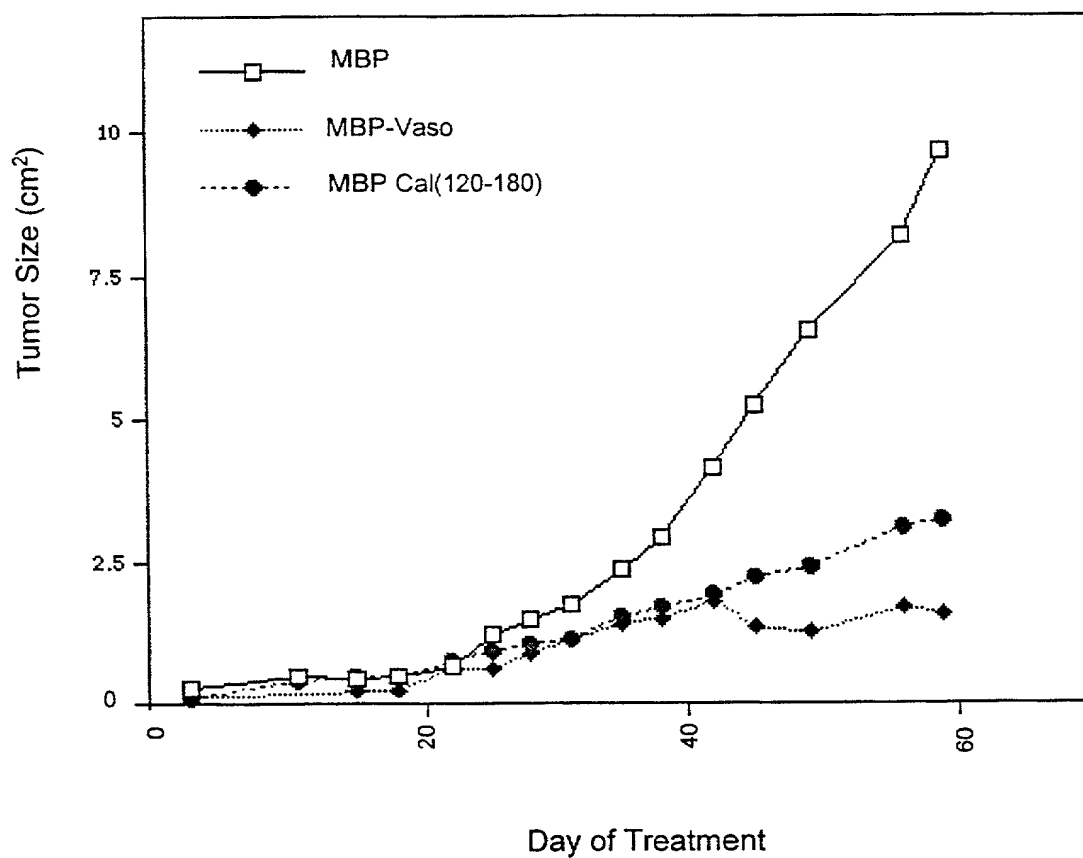


Fig. 16

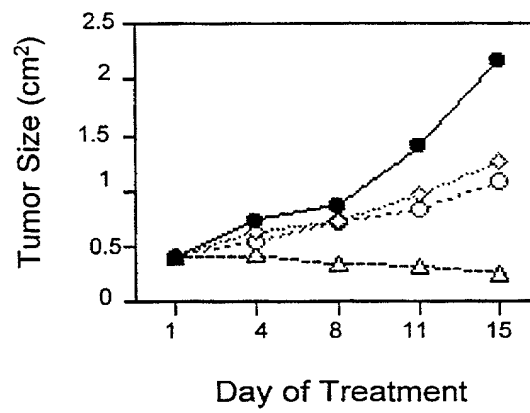


Fig. 17A

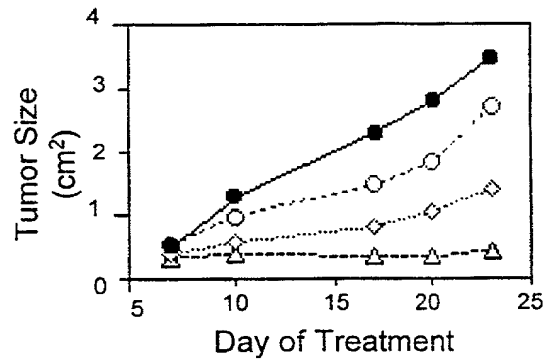


Fig. 17B

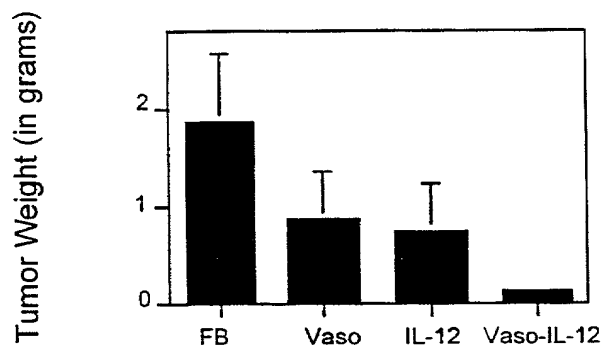


Fig. 17C

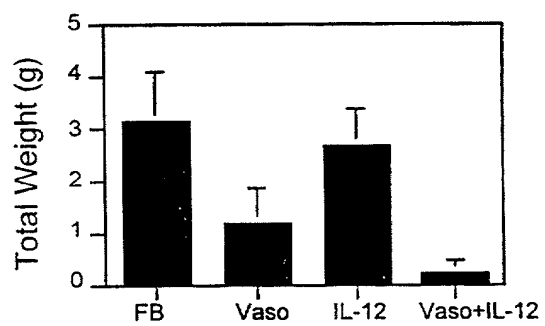


Fig. 17D

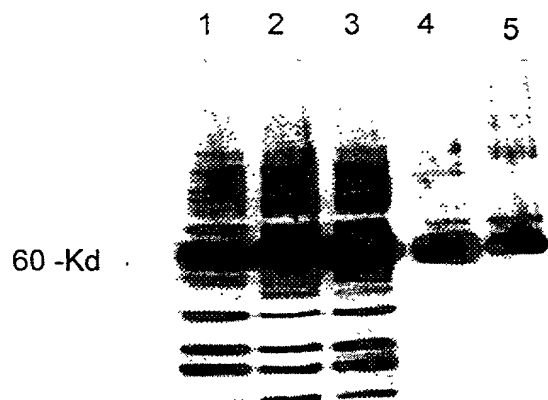


Fig. 18

4/04/01 WED 16:01 FAX 226 9448

WDN/SAS:jlb 4239-58378 E-082-98/2 04/04/01

**COMBINED DECLARATION AND POWER OF ATTORNEY
FOR PATENT APPLICATION**

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled **USE OF CALRETICULIN AND CALRETICULIN FRAGMENTS TO INHIBIT ENDOTHELIAL CELL GROWTH AND ANGIOGENESIS, AND SUPPRESS TUMOR GROWTH**, the specification of which

- ☒ is attached hereto.
- ☐ was filed on _____ as United States Application No. _____.
- ☒ was filed on October 5, 1999 as International Application No. PCT/US99/23240.
- ☐ and was amended on _____ (if applicable).
- ☐ with amendments through _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56. If this is a continuation-in-part application filed under the conditions specified in 35 U.S.C. § 120 which discloses and claims subject matter in addition to that disclosed in the prior copending application, I further acknowledge the duty to disclose material information as defined in 37 CFR § 1.56 which occurred between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under Title 35, United States Code, § 119(a)-(d) of any foreign application(s) for patent or inventor's certificate or of an PCT International application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign application(s) for patent or inventor's certificate or any PCT International application(s) designating at least one country other than the United States of America filed by me on the same subject matter having a filing date before that of the application(s) on which priority is claimed:

Prior Foreign Application(s)	Priority Claimed
	<input type="checkbox"/> Yes <input type="checkbox"/> No

I hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States provisional application(s) listed below:

60/103,438	October 6, 1998
(Application No.)	(Filing Date)

Apr 06 01 12:09p

NCI

301-594-9590

p. 3
003

KLARQUIST SPARKMAN

4/04/01 WED 16:02 FAX 228 9446
WDN/SAS:jib 4239-58378 E-082-98/2 04/04/01

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) or § 365(c) of any PCT International application(s) designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 1.56(a) which occurred between the filing date of the prior application and the national or PCT International filing date of this application:

PCT/US99/23240	October 5, 1999	Pending
(Application No.)	(Filing Date)	(Status: patented, pending, abandoned)

The undersigned hereby authorizes the U.S. attorney or agent named herein to accept and follow instructions from _____ as to any action to be taken in the Patent and Trademark Office regarding this application without direct communication between the U.S. attorney or agent and the undersigned. In the event of a change in the persons from whom instructions may be taken, the U.S. attorney or agent named herein will be so notified by the undersigned.

I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application, to file a corresponding international application, and to transact all business in the Patent and Trademark Office connected therewith:

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John Peter Kim	<u>38,514</u>	Norbert J. Pontzer	<u>40,777</u>
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Stacey C. Slater	<u>36,011</u>	Gregory L. Maurer	<u>43,781</u>
Susan Alpert Siegel	<u>43,121</u>	Robert F. Scotti	<u>39,830</u>
Wayne W. Rupert	<u>34,420</u>	Deborah B. Blyveis	<u>47,337</u>

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Inventor's Signature

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Date

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Inventor's Signature

Sandra E Pike

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Date

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April 5, 2001
Date

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MD

Citizenship: United States of America

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SEQUENCE LISTING

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ccttcttgga gccagagga gggcagcaga aggggggtggt gtctccaacc cccagcact 1782

gaggaagaac ggggctcttc tcatttcacc cctccctttc tcccctgccc ccaggactgg 1842

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<213> Homo sapiens

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<213> Homo sapiens

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Asn Val Leu Ile Asn Lys Asp Ile Arg Cys Lys Asp Asp Glu Phe Thr
 20 25 30

His Leu Tyr Thr Leu Ile Val Arg Pro Asp Asn Thr Tyr Glu Val Lys
 35 40 45

Ile Asp Asn Ser Gln Val Glu Ser Gly Ser Leu Glu Asp Asp Trp Asp
 50 55 60

Phe Leu Pro Pro Lys Lys Ile Lys Asp Pro Asp Ala Ser Lys Pro Glu
 65 70 75 80

Asp Trp Asp Glu Arg Ala Lys Ile Asp Asp Pro Thr Asp Ser Lys Pro
 85 90 95

Glu Asp Trp Asp Lys Pro Glu His Ile Pro Asp Pro Asp Ala Lys Lys
 100 105 110

Pro Glu Asp Trp Asp Glu Glu Met Asp Gly Glu Trp Glu Pro Pro Val
 115 120 125

Ile Gln Asn Pro Glu Tyr Lys Gly Glu Trp Lys Pro Arg Gln Ile Asp
 130 135 140

Asn Pro Asp Tyr Lys Gly Thr Trp Ile His Pro Glu Ile Asp Asn Pro
 145 150 155 160

Glu Tyr Ser Pro Asp Pro Ser Ile Tyr Ala Tyr Asp Asn Phe Gly Val
 165 170 175

Leu Gly Leu Asp Leu Trp Gln Val Lys Ser Gly Thr Ile Phe Asp Asn
 180 185 190

Phe Leu Ile Thr Asn Asp Glu Ala Tyr Ala Glu Glu Phe Gly Asn Glu
 195 200 205

Thr Trp Gly Val Thr Lys Ala Ala Glu Lys Gln Met Lys Asp Lys Gln
 210 215 220

Asp Glu Glu Gln Arg Leu Lys Glu Glu Glu Glu Asp Lys Lys Arg Lys
 225 230 235 240

Glu Glu Glu Glu Ala Glu Asp Lys Glu Asp Asp Glu Asp Lys Asp Glu
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Pro Gly Gln Ala Lys Asp Glu Leu

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<210> 12
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<210> 18

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<210> 21

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<400> 29

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RAW SEQUENCE LISTING

DATE: 07/23/2001

PATENT APPLICATION: US/09/807,148

TIME: 18:05:42

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Output Set: N:\CRF3\07232001\I807148.raw

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3 <110> APPLICANT: Tosato, Giovanna et al.
5 <120> TITLE OF INVENTION: Use of Calreticulin and Calretuculin Fragments to
6 Inhibit Endothelial Cell Growth and Angiogenesis, and
7 Suppress Tumor Growth
9 <130> FILE REFERENCE: 4239 53372
C--> 11 <140> CURRENT APPLICATION NUMBER: US/09/807,148
C--> 12 <141> CURRENT FILING DATE: 2001-06-04
14 <150> PRIOR APPLICATION NUMBER: US 60/103,438
15 <151> PRIOR FILING DATE: 1998-10-06
17 <160> NUMBER OF SEQ ID NOS: 35
19 <170> SOFTWARE: PatentIn Ver. 2.0
21 <210> SEQ ID NO: 1
22 <211> LENGTH: 1251
23 <212> TYPE: DNA
24 <213> ORGANISM: Homo sapiens
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27 <221> NAME/KEY: CDS
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36	Ala	Glu	Pro	Ala	Val	Tyr	Phe	Lys	Glu	Gln	Phe	Leu	Asp	Gly	Asp	Gly	
37					20				25					30			
39	tgg	act	tcc	cgc	tgg	atc	gaa	tcc	aaa	cac	aag	tca	gat	ttt	ggc	aaa	144
40	Trp	Thr	Ser	Arg	Trp	Ile	Glu	Ser	Lys	His	Lys	Ser	Asp	Phe	Gly	Lys	
41					35				40					45			
43	ttc	gtt	ctc	agt	tcc	ggc	aag	ttc	tac	ggt	gac	gag	gag	aaa	gat	aaa	192
44	Phe	Val	Leu	Ser	Ser	Gly	Lys	Phe	Tyr	Gly	Asp	Glu	Glu	Lys	Asp	Lys	
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47	ggt	ttg	cag	aca	agc	cag	gat	gca	cgc	ttt	tat	gct	ctg	tcg	gcc	agt	240
48	Gly	Leu	Gln	Thr	Ser	Gln	Asp	Ala	Arg	Phe	Tyr	Ala	Leu	Ser	Ala	Ser	
49	65					70				75					80		
51	ttc	gag	cct	ttc	agc	aac	aaa	ggc	cag	acg	ctg	gtg	gtg	cag	ttc	acg	288
52	Phe	Glu	Pro	Phe	Ser	Asn	Lys	Gly	Gln	Thr	Leu	Val	Val	Gln	Phe	Thr	
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55	gtg	aaa	cat	gag	cag	aac	atc	gac	tgt	ggg	ggc	ggc	tat	gtg	aag	ctg	336
56	Val	Lys	His	Glu	Gln	Asn	Ile	Asp	Cys	Gly	Gly	Gly	Tyr	Val	Lys	Leu	
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60	Phe	Pro	Asn	Ser	Leu	Asp	Gln	Thr	Asp	Met	His	Gly	Asp	Ser	Glu	Tyr	
61					115				120					125			
63	aac	atc	atg	ttt	ggt	ccc	gac	atc	tgt	ggc	cct	ggc	acc	aag	aag	gtt	432
64	Asn	Ile	Met	Phe	Gly	Pro	Asp	Ile	Cys	Gly	Pro	Gly	Thr	Lys	Lys	Val	
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RAW SEQUENCE LISTING

PATENT APPLICATION: US/09/807,148

DATE: 07/23/2001

TIME: 18:05:42

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72 Ile Arg Cys Lys Asp Asp Glu Phe Thr His Leu Tyr Thr Leu Ile Val
73 165 170 175
75 cgg cca gac aac acc tat gag gtg aag att gac aac agc cag gtg gag 576
76 Arg Pro Asp Asn Thr Tyr Glu Val Lys Ile Asp Asn Ser Gln Val Glu
77 180 185 190
79 tcc ggc tcc ttg gaa gac gat tgg gac ttc ctg cca ccc aag aag ata 624
80 Ser Gly Ser Leu Glu Asp Asp Trp Asp Phe Leu Pro Pro Lys Lys Ile
81 195 200 205
83 aag gat cct gat gct tca aaa ccg gaa gac tgg gat gag cgg gcc aag 672
84 Lys Asp Pro Asp Ala Ser Lys Pro Glu Asp Trp Asp Glu Arg Ala Lys
85 210 215 220
87 atc gat gat ccc aca gac tcc aag cct gag gac tgg gac aag ccc gag 720
88 Ile Asp Asp Pro Thr Asp Ser Lys Pro Glu Asp Trp Asp Lys Pro Glu
89 225 230 235 240
91 cat atc cct gac cct gat gct aag aag ccc gag gac tgg gat gaa gag 768
92 His Ile Pro Asp Pro Asp Ala Lys Lys Pro Glu Asp Trp Asp Glu Glu
93 245 250 255
95 atg gac gga gag tgg gaa ccc cca gtg att cag aac cct gag tac aag 816
96 Met Asp Gly Glu Trp Glu Pro Pro Val Ile Gln Asn Pro Glu Tyr Lys
97 260 265 270
99 ggt gag tgg aag ccc cgg cag atc gac aac cca gat tac aag ggc act 864
100 Gly Glu Trp Lys Pro Arg Gln Ile Asp Asn Pro Asp Tyr Lys Gly Thr
101 275 280 285
103 tgg atc cac cca gaa att gac aac ccc gag tat tct ccc gat ccc agt 912
104 Trp Ile His Pro Glu Ile Asp Asn Pro Glu Tyr Ser Pro Asp Pro Ser
105 290 295 300
107 atc tat gcc tat gat aac ttt ggc gtg ctg ggc ctg gac ctc tgg cag 960
108 Ile Tyr Ala Tyr Asp Asn Phe Gly Val Leu Gly Leu Asp Leu Trp Gln
109 305 310 315 320
111 gtc aag tct ggc acc atc ttt gac aac ttc ctc atc acc aac gat gag 1008
112 Val Lys Ser Gly Thr Ile Phe Asp Asn Phe Leu Ile Thr Asn Asp Glu
113 325 330 335
115 gca tac gct gag gag ttt ggc aac gag acg tgg ggc gta aca aag gca 1056
116 Ala Tyr Ala Glu Glu Phe Gly Asn Glu Thr Trp Gly Val Thr Lys Ala
117 340 345 350
119 gca gag aaa caa atg aag gac aaa cag gac gag gag cag agg ctt aag 1104
120 Ala Glu Lys Gln Met Lys Asp Lys Gln Asp Glu Glu Gln Arg Leu Lys
121 355 360 365
123 gag gag gaa gaa gac aag aaa cgc aaa gag gag gag gag gca gag gac 1152
124 Glu Glu Glu Glu Asp Lys Lys Arg Lys Glu Glu Glu Glu Ala Glu Asp
125 370 375 380
127 aag gag gat gat gag gac aaa gat gag gat gag gag gat gag gag gac 1200
128 Lys Glu Asp Asp Glu Asp Lys Asp Glu Asp Glu Glu Asp Glu Glu Asp
129 385 390 395 400
131 aag gag gaa gat gag gag gaa gat gtc ccc ggc cag gcc aag gac gag 1248
132 Lys Glu Glu Asp Glu Glu Glu Asp Val Pro Gly Gln Ala Lys Asp Glu

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RAW SEQUENCE LISTING

PATENT APPLICATION: US/09/807,148

DATE: 07/23/2001

TIME: 18:05:42

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Output Set: N:\CRF3\07232001\I807148.raw

133 405 410 415 1251

135 ctg

136 Leu

139 <210> SEQ ID NO: 2

140 <211> LENGTH: 417

141 <212> TYPE: PRT

142 <213> ORGANISM: Homo sapiens

144 <400> SEQUENCE: 2

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146 1 5 10 15

148 Ala Glu Pro Ala Val Tyr Phe Lys Glu Gln Phe Leu Asp Gly Asp Gly

149 20 25 30

151 Trp Thr Ser Arg Trp Ile Glu Ser Lys His Lys Ser Asp Phe Gly Lys

152 35 40 45

154 Phe Val Leu Ser Ser Gly Lys Phe Tyr Gly Asp Glu Glu Lys Asp Lys

155 50 55 60

157 Gly Leu Gln Thr Ser Gln Asp Ala Arg Phe Tyr Ala Leu Ser Ala Ser

158 65 70 75 80

160 Phe Glu Pro Phe Ser Asn Lys Gly Gln Thr Leu Val Val Gln Phe Thr

161 85 90 95

163 Val Lys His Glu Gln Asn Ile Asp Cys Gly Gly Gly Tyr Val Lys Leu

164 100 105 110

166 Phe Pro Asn Ser Leu Asp Gln Thr Asp Met His Gly Asp Ser Glu Tyr

167 115 120 125

169 Asn Ile Met Phe Gly Pro Asp Ile Cys Gly Pro Gly Thr Lys Lys Val

170 130 135 140

172 His Val Ile Phe Asn Tyr Lys Gly Lys Asn Val Leu Ile Asn Lys Asp

173 145 150 155 160

175 Ile Arg Cys Lys Asp Asp Glu Phe Thr His Leu Tyr Thr Leu Ile Val

176 165 170 175

178 Arg Pro Asp Asn Thr Tyr Glu Val Lys Ile Asp Asn Ser Gln Val Glu

179 180 185 190

181 Ser Gly Ser Leu Glu Asp Asp Trp Asp Phe Leu Pro Pro Lys Lys Ile

182 195 200 205

184 Lys Asp Pro Asp Ala Ser Lys Pro Glu Asp Trp Asp Glu Arg Ala Lys

185 210 215 220

187 Ile Asp Asp Pro Thr Asp Ser Lys Pro Glu Asp Trp Asp Lys Pro Glu

188 225 230 235 240

190 His Ile Pro Asp Pro Asp Ala Lys Lys Pro Glu Asp Trp Asp Glu Glu

191 245 250 255

193 Met Asp Gly Glu Trp Glu Pro Pro Val Ile Gln Asn Pro Glu Tyr Lys

194 260 265 270

196 Gly Glu Trp Lys Pro Arg Gln Ile Asp Asn Pro Asp Tyr Lys Gly Thr

197 275 280 285

199 Trp Ile His Pro Glu Ile Asp Asn Pro Glu Tyr Ser Pro Asp Pro Ser

200 290 295 300

202 Ile Tyr Ala Tyr Asp Asn Phe Gly Val Leu Gly Leu Asp Leu Trp Gln

203 305 310 315 320

205 Val Lys Ser Gly Thr Ile Phe Asp Asn Phe Leu Ile Thr Asn Asp Glu

RAW SEQUENCE LISTING

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208 Ala Tyr Ala Glu Glu Phe Gly Asn Glu Thr Trp Gly Val Thr Lys Ala
209          340          345          350
211 Ala Glu Lys Gln Met Lys Asp Lys Gln Asp Glu Glu Gln Arg Leu Lys
212          355          360          365
214 Glu Glu Glu Glu Asp Lys Lys Arg Lys Glu Glu Glu Glu Ala Glu Asp
215          370          375          380
217 Lys Glu Asp Asp Glu Asp Lys Asp Glu Asp Glu Glu Asp Glu Glu Asp
218 385          390          395          400
220 Lys Glu Glu Asp Glu Glu Glu Asp Val Pro Gly Gln Ala Lys Asp Glu
221          405          410          415
223 Leu
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228 <211> LENGTH: 400
229 <212> TYPE: PRT
230 <213> ORGANISM: Homo sapiens
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237          20          25          30
239 Val Leu Ser Ser Gly Lys Phe Tyr Gly Asp Glu Glu Lys Asp Lys Gly
240          35          40          45
242 Leu Gln Thr Ser Gln Asp Ala Arg Phe Tyr Ala Leu Ser Ala Ser Phe
243          50          55          60
245 Glu Pro Phe Ser Asn Lys Gly Gln Thr Leu Val Val Gln Phe Thr Val
246 65          70          75          80
248 Lys His Glu Gln Asn Ile Asp Cys Gly Gly Tyr Val Lys Leu Phe
249          85          90          95
251 Pro Asn Ser Leu Asp Gln Thr Asp Met His Gly Asp Ser Glu Tyr Asn
252          100          105          110
254 Ile Met Phe Gly Pro Asp Ile Cys Gly Pro Gly Thr Lys Lys Val His
255          115          120          125
257 Val Ile Phe Asn Tyr Lys Gly Lys Asn Val Leu Ile Asn Lys Asp Ile
258          130          135          140
260 Arg Cys Lys Asp Asp Glu Phe Thr His Leu Tyr Thr Leu Ile Val Arg
261 145          150          155          160
263 Pro Asp Asn Thr Tyr Glu Val Lys Ile Asp Asn Ser Gln Val Glu Ser
264          165          170          175
266 Gly Ser Leu Glu Asp Asp Trp Asp Phe Leu Pro Pro Lys Lys Ile Lys
267          180          185          190
269 Asp Pro Asp Ala Ser Lys Pro Glu Asp Trp Asp Glu Arg Ala Lys Ile
270          195          200          205
272 Asp Asp Pro Thr Asp Ser Lys Pro Glu Asp Trp Asp Lys Pro Glu His
273          210          215          220
275 Ile Pro Asp Pro Asp Ala Lys Lys Pro Glu Asp Trp Asp Glu Glu Met
276 225          230          235          240
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RAW SEQUENCE LISTING

PATENT APPLICATION: US/09/807,148

DATE: 07/23/2001

TIME: 18:05:42

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281 Glu Trp Lys Pro Arg Gln Ile Asp Asn Pro Asp Tyr Lys Gly Thr Trp
 282 260 265 270
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 285 275 280 285
 287 Tyr Ala Tyr Asp Asn Phe Gly Val Leu Gly Leu Asp Leu Trp Gln Val
 288 290 295 300
 290 Lys Ser Gly Thr Ile Phe Asp Asn Phe Leu Ile Thr Asn Asp Glu Ala
 291 305 310 315 320
 293 Tyr Ala Glu Glu Phe Gly Asn Glu Thr Trp Gly Val Thr Lys Ala Ala
 294 325 330 335
 296 Glu Lys Gln Met Lys Asp Lys Gln Asp Glu Glu Gln Arg Leu Lys Glu
 297 340 345 350
 299 Glu Glu Glu Asp Lys Lys Arg Lys Glu Glu Glu Glu Ala Glu Asp Lys
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 306 385 390 395 400
 312 <210> SEQ ID NO: 4
 313 <211> LENGTH: 180
 314 <212> TYPE: PRT
 315 <213> ORGANISM: Homo sapiens
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 324 Val Leu Ser Ser Gly Lys Phe Tyr Gly Asp Glu Glu Lys Asp Lys Gly
 325 35 40 45
 327 Leu Gln Thr Ser Gln Asp Ala Arg Phe Tyr Ala Leu Ser Ala Ser Phe
 328 50 55 60
 330 Glu Pro Phe Ser Asn Lys Gly Gln Thr Leu Val Val Gln Phe Thr Val
 331 65 70 75 80
 333 Lys His Glu Gln Asn Ile Asp Cys Gly Gly Gly Tyr Val Lys Leu Phe
 334 85 90 95
 336 Pro Asn Ser Leu Asp Gln Thr Asp Met His Gly Asp Ser Glu Tyr Asn
 337 100 105 110
 339 Ile Met Phe Gly Pro Asp Ile Cys Gly Pro Gly Thr Lys Lys Val His
 340 115 120 125
 342 Val Ile Phe Asn Tyr Lys Gly Lys Asn Val Leu Ile Asn Lys Asp Ile
 343 130 135 140
 345 Arg Cys Lys Asp Asp Glu Phe Thr His Leu Tyr Thr Leu Ile Val Arg
 346 145 150 155 160
 348 Pro Asp Asn Thr Tyr Glu Val Lys Ile Asp Asn Ser Gln Val Glu Ser
 349 165 170 175
 351 Gly Ser Leu Glu
 352 180
 355 <210> SEQ ID NO: 5
 356 <211> LENGTH: 61

Use of n and/or Xaa has been detected in the Sequence Listing.
 Review the Sequence Listing to insure a corresponding
 explanation is presented in the <220> to <223> fields of
 each sequence using n or Xaa.

VERIFICATION SUMMARY

PATENT APPLICATION: US/09/807,148

DATE: 07/23/2001

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FOUO - 07/23/01

PCT

RAW SEQUENCE LISTING
 PATENT APPLICATION: US/09/807,148

DATE: 07/05/2001
 TIME: 15:36:34

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Does Not Comply
 Corrected Diskette Needed

3 <110> APPLICANT: Tosato, Giovanna et al.
 5 <120> TITLE OF INVENTION: Use of Calreticulin and Calretuculin Fragments to
 6 Inhibit Endothelial Cell Growth and Angiogenesis, and
 7 Suppress Tumor Growth
 9 <130> FILE REFERENCE: 4239 53372
 C--> 11 <140> CURRENT APPLICATION NUMBER: US/09/807,148
 C--> 12 <141> CURRENT FILING DATE: 2001-06-04
 14 <150> PRIOR APPLICATION NUMBER: US 60/103,438
 15 <151> PRIOR FILING DATE: 1998-10-06
 17 <160> NUMBER OF SEQ ID NOS: 35
 19 <170> SOFTWARE: PatentIn Ver. 2.0

ERRORED SEQUENCES

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VERIFICATION SUMMARY

DATE: 07/05/2001

PATENT APPLICATION: US/09/807,148

TIME: 15:36:36

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T03040"04F20350

PTO/PCT Rec'd 04 JUN 2001

PATENT #3

Attorney's Matter No. 4239-58378

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of: Tosato et al.

Art Unit: To be assigned

Application No. 09/807,148

CERTIFICATE OF MAILING

Filed: April 5, 2001

I hereby certify that this paper and the documents referred to as being attached or enclosed herewith are being deposited with the United States Postal Service on May 29, 2001 as First Class Mail in an envelope addressed to: BOX PCT, COMMISSIONER FOR PATENTS, WASHINGTON D C. 20231.

For: USE OF CALRETICULIN AND
CALRETICULIN FRAGMENTS TO
INHIBIT ENDOTHELIAL CELL
GROWTH AND ANGIOGENESIS, AND
SUPPRESS TUMOR GROWTH



William D. Noonan, M.D.
Attorney for Applicant

Examiner: To be assigned

Date: May 29, 2001

STATEMENT IN COMPLIANCE WITH 37 C.F.R. § 1.821(f)

BOX PCT
COMMISSIONER FOR PATENTS
Washington, DC 20231

Sir:

In compliance with 37 C.F.R. § 1.821(f), the undersigned declares that the nucleotide and/or amino acid sequences presented in the paper copy of the "Sequence Listing" submitted herewith are the same as the sequences contained in the computer-readable form of said "Sequence Listing." No new matter has been added.

Respectfully submitted,

KLARQUIST SPARKMAN CAMPBELL
LEIGH & WHINSTON, LLP

By 

William D. Noonan, M.D.
Registration No. 30,878

One World Trade Center, Suite 1600
121 S.W. Salmon Street
Portland, Oregon 97204
Telephone: (503) 226-7391
Facsimile: (503) 228-9446

09/807148-04501

SEQUENCE LISTING

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 Ile Asp Asn Ser Gln Val Glu Ser Gly Ser Leu Glu Asp Asp Trp Asp
 50 55 60

Phe Leu Pro Pro Lys Lys Ile Lys Asp Pro Asp Ala Ser Lys Pro Glu
65 70 75 80

Asp Trp Asp Glu Arg Ala Lys Ile Asp Asp Pro Thr Asp Ser Lys Pro
85 90 95

Glu Asp Trp Asp Lys Pro Glu His Ile Pro Asp Pro Asp Ala Lys Lys
100 105 110

Pro Glu Asp Trp Asp Glu Glu Met Asp Gly Glu Trp Glu Pro Pro Val
115 120 125

Ile Gln Asn Pro Glu Tyr Lys Gly Glu Trp Lys Pro Arg Gln Ile Asp
130 135 140

Asn Pro Asp Tyr Lys Gly Thr Trp Ile His Pro Glu Ile Asp Asn Pro
145 150 155 160

Glu Tyr Ser Pro Asp Pro Ser Ile Tyr Ala Tyr Asp Asn Phe Gly Val
165 170 175

Leu Gly Leu Asp Leu Trp Gln Val Lys Ser Gly Thr Ile Phe Asp Asn
180 185 190

Phe Leu Ile Thr Asn Asp Glu Ala Tyr Ala Glu Glu Phe Gly Asn Glu
195 200 205

Thr Trp Gly Val Thr Lys Ala Ala Glu Lys Gln Met Lys Asp Lys Gln
210 215 220

Asp Glu Glu Gln Arg Leu Lys Glu Glu Glu Glu Asp Lys Lys Arg Lys
225 230 235 240

Glu Glu Glu Glu Ala Glu Asp Lys Glu Asp Asp Glu Asp Lys Asp Glu
245 250 255

Asp Glu Glu Asp Glu Glu Asp Lys Glu Glu Asp Glu Glu Glu Asp Val
260 265 270

Pro Gly Gln Ala Lys Asp Glu Leu
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<220>
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<220>
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integrin sequence

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<223> Description of Artificial Sequence:Consensus
steroid nuclear receptor sequence

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<210> 12
<211> 7
<212> PRT
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<220>
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<210> 15
<211> 7
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<210> 16
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 steroid nuclear receptor

<400> 21
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steroid nuclear receptor

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<210> 23
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<220>
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steroid nuclear receptor

<400> 23
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steroid nuclear receptor

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<223> Description of Artificial Sequence:Portion of steroid nuclear receptor

Thr Cys Gly Ser Cys Lys Val Phe Phe Lys Arg Ala Met Glu Gly
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<223> Description of Artificial Sequence:Portion of steroid nuclear receptor

Ser Cys Glu Gly Cys Lys Ala Phe Phe Lys Arg Ser Ile Gln Gly
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<223> Description of Artificial Sequence:Portion of steroid nuclear receptor

Ser Cys Glu Gly Cys Lys Gly Phe Phe Lys Arg Thr Val Arg Lys
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Thr Cys Glu Gly Cys Thr Gly Phe Phe Lys Arg Ser Ile Arg Lys
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17

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